

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
5 June 2003 (05.06.2003)

PCT

(10) International Publication Number
WO 03/045434 A2

(51) International Patent Classification⁷: **A61K 45/06**,
38/12, 31/485, A61P 25/04 // (A61K 38/12, 31:485)

(21) International Application Number: PCT/US02/37461

(22) International Filing Date:
22 November 2002 (22.11.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/333,599 27 November 2001 (27.11.2001) US

(71) Applicant: **THE BOARD OF TRUSTEES OF THE
UNIVERSITY OF ILLINOIS** [US/US]; 352 Administra-
tion Building, 506 Wright Street, Urbana, IL 61801 (US).

(72) Inventor: **GULATI, Anil**; 1929 Lisson Road, Naperville,
IL 60565 (US).

(74) Agent: **NAPOLI, James, J.**; Marshall, Gerstein & Borun,
6300 Sears Tower, 233 South Wacker Drive, Chicago, IL
60606 (US).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VC,
VN, YU, ZA, ZM, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK,
TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG).

Published:

— without international search report and to be republished
upon receipt of that report

*For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.*

(54) Title: METHOD AND COMPOSITION FOR POTENTIATING AN OPIATE ANALGESIC

(57) Abstract: Composition and methods of treating pain and reducing or reversing tolerance to opiate analgesics are disclosed. The composition and method utilize an opiate analgesic and an endothelin antagonist as active agents to treat pain in mammals, including humans.



WO 03/045434 A2

- 1 -

**METHOD AND COMPOSITION FOR POTENTIATING
AN OPIATE ANALGESIC**

CROSS REFERENCE TO RELATED APPLICATIONS

This application claims the benefit of
5 U.S. provisional patent application Serial No.
60/333,599, filed November 27, 2001.

FIELD OF THE INVENTION

The present invention relates to the
treatment of pain using an opiate analgesic and an
10 endothelin receptor antagonist. More particularly,
the present invention relates to a method of poten-
tiating the effects of an opiate analgesic, like
morphine, in a mammal by administration of a thera-
peutically effective amount of an endothelin re-
15 ceptor antagonist. The composition and method
permit a reduction in the opiate analgesic dose to
provide a desired analgesic effect, without effect-
ing the cataleptic action of the opiate analgesic.
The present invention also relates to a method of
20 reducing or reversing tolerance to an opiate anal-
gesic in an individual undergoing opiate analgesic
treatment by administering a therapeutically effec-
tive amount of an endothelin receptor antagonist.
The present composition and methods also reduce the
25 incidence of opiate analgesic addiction.

- 2 -

BACKGROUND OF THE INVENTION

Analgesics are agents that relieve pain by acting centrally to elevate pain threshold, preferably without disturbing consciousness or altering
5 other sensory functions. A mechanism by which analgesic drugs obtund pain (i.e., raise the pain threshold) has been formulated. Research in this area has resulted in the development of a number of opiate and opioid analgesics having diverse pharmacological actions.
10

The available opiate and opioid analgesics are derivatives of five chemical groups (i.e., phenanthrenes, phenylheptylamines, phenylpiperidines, morphinans, and benzomorphans). Pharmacologically,
15 these opiates and nonopiates differ significantly in activity. Some are strong agonists (morphine), others are moderates-to-mild agonists (codeine). In contrast, some opiate derivatives exhibit mixed agonist-antagonist activity (nalbuphine), whereas
20 others are opiate antagonists (naloxone). Morphine is the prototype of the opiate and opioid analgesics, all of which have similar actions on the central nervous system.

Morphine is an alkaloid chemically derived
25 from opium *papaver somniferum*. Other drugs, such as heroin, are processed from morphine or codeine. Such opiates have been used both medically and non-medically for centuries. By the early 19th century, morphine had been extracted in a pure form suitable
30 for solution. With the introduction of the hypodermic needle, injection of a morphine solution

- 3 -

became the common method of administration. Of the twenty alkaloids contained in opium, only codeine and morphine are still in widespread clinical use.

The opiates are among the most powerfully
5 acting and clinically useful drugs producing depression of the central nervous system. Drugs of this group are used principally as analgesics, but possess numerous other useful properties. Morphine, for example, is used to relieve pain, induce sleep
10 in the presence of pain, check diarrhea, suppress cough, ease dyspnea, and facilitate anesthesia.

However, morphine also depresses respiration; increases the activity and tone of the smooth muscles of the gastrointestinal, biliary, and
15 urinary tracts causing constipation, gallbladder spasm, and urinary retention; causes nausea and vomiting in some individuals; and can induce cutaneous pruritus. In addition, morphine and related compounds have other properties that tend to limit
20 their usefulness.

For example, when morphine and related compounds are administered over a long time period, tolerance to the analgesic effect develops, and the dose then must be increased periodically to obtain
25 equivalent pain relief. Eventually, tolerance and physical dependence develop, which, combined with euphoria, result in excessive use and addiction of those patients having susceptible personalities. For these reasons, morphine and its derivatives must
30 be used only as directed by a physician (i.e., not in greater dose, more often, or longer than pre-

- 4 -

scribed), and should not be used to treat pain when a different analgesic will suffice.

Nevertheless, morphine remains the major drug for the treatment of moderate to severe pain (Foley, 1993). Opioids particularly are used to
5 treat conditions lacking a standard treatment, such as cancer pain, trauma, myocardial infarction, post-operative pain, and neuropathic pain. However, opioid painkillers have significant adverse side
10 effects like respiratory depression, nausea, vomiting, dizziness, sedation, mental clouding, constipation, urinary retention, and severe itching.

These adverse side effects limit the usefulness of opioids, like morphine, as painkillers.
15 Therefore, several companies are developing a new generation of opioid painkillers, but advances in neuroscience have not progressed a sufficient extent to provide a significant breakthrough. Typically, companies are using proprietary technology to re-
20 formulate opioid drugs, such as morphine, into branded painkillers with improved clinical benefits. To date, innovations in the field of opioid painkillers have largely focused on increasing the convenience of opioid drugs. For example, important
25 advances have been made in opioid delivery, such as sustained release formulations and transmucosal delivery.

The present invention is directed to the discovery that some pharmacological actions of morphine can be modified by coadministration of an
30 endothelin receptor antagonist, hereafter termed an

- 5 -

"endothelin antagonist." U.S. patent publication US 2002/0082285 A1 discloses the use of an endothelin antagonist in the treatment of pain.

SUMMARY OF THE INVENTION

5 The present invention is directed to administration of an endothelin antagonist in combination with an opiate analgesic. More particularly, administration of an opiate analgesic in combination with an endothelin antagonist potentiates the anal-
10 gesic effect of opioids, and, therefore, lowers the dose of analgesic required to provide a desired pain-reducing effect, without affecting the cataleptic properties of the analgesic. The reduced amount of opiate analgesic required to provide a
15 desired effect reduces the severity of various adverse side effects associated with opiate analgesic treatment.

 Accordingly, one aspect of the present invention is to provide a composition comprising an
20 opiate analgesic, e.g., morphine, and an endothelin antagonist for use in treating pain. Such a composition provides a safety factor for the patient because endothelin significantly regulates the autonomic nervous system (A. Gulati et al., (1997); A.
25 Kumar et al. (1997)), and a majority of the withdrawal reactions of morphine also are mediated through the autonomic nervous system. Therefore, endothelin is expected to modulate various pharmacological actions of morphine and other opiate
30 analgesics.

- 6 -

The present invention also is directed to a method of reducing or reversing tolerance to an opiate analgesic in an individual undergoing an opiate analgesic therapy by administering an
5 endothelin antagonist to the individual. In the absence of an administered dose of endothelin antagonist, the opiate analgesic dose would have to be increased over time to achieve the same pain-reducing effect. Administration of an endothelin
10 antagonist allows the opiate analgesic to be administered at a constant, or reduced, dose to achieve a desired pain treatment. The constant or reduced amount of opiate analgesic required to provide a desired pain-reducing effect thus reduces the
15 severity of various adverse side effects associated with opiate analgesic treatment, and reduces the possibility of opiate analgesic dependence.

The present invention also provides a method for improved pain treatment. In particular,
20 the present invention is directed to methods of using an opiate analgesic and an endothelin antagonist to prevent and/or treat pain. More particularly, the present invention is directed to compositions containing morphine and an endothelin
25 antagonist, and to use of an opiate analgesic and endothelin antagonist, administered simultaneously or sequentially, in methods of treating pain and reducing or reversing opiate analgesic tolerance and dependence.

30 An important aspect of the present invention, therefore, is to provide a method and composi-

- 7 -

tion for preventing or treating pain, while reducing the occurrence or severity of adverse side effects associated with opiate analgesic treatment.

Another aspect of the present invention is to reduce the problem of dependence and addiction associated with present opiate analgesics used to treat pain.

Still another aspect of the present invention is to provide a method of reducing or reversing opiate analgesic tolerance in an individual undergoing an opiate analgesic therapy by administering a therapeutically effective amount of an endothelin antagonist to the individual.

Yet another aspect of the present invention is to provide an article of manufacture for human pharmaceutical use, comprising (a) a package insert, (b) a container, and either (c1) a packaged composition comprising an opiate analgesic and an endothelin antagonist or (c2) a packaged composition comprising an opiate analgesic and a packaged composition comprising an endothelin antagonist.

These and other aspects of the present invention will become apparent from the following detailed description of the preferred embodiments of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 illustrates an example of the groups of rats treated with a vehicle, endothelin antagonist, morphine, and endothelin antagonist plus morphine;

- 8 -

Fig. 2 contains plots for the effect of BQ123 pretreatment on analgesia induced by morphine for four groups of treated rats;

5 Figs. 3-5 contain plots for the effect of BQ123 pretreatment on analgesia induced by morphine (2, 4, and 8 mg/kg, respectively), for four groups of treated mice; and

Figs. 6-8 contain plots showing a reduced morphine tolerance in mice after treatment with the
10 endothelin antagonists BQ123 or BMS182874.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention is directed to the simultaneous or sequential administration of an opiate analgesic and an endothelin antagonist to
15 prevent and/or treat pain. In particular, the administration of morphine and an endothelin antagonist to rats and mice show that the endothelin antagonist potentiates the analgesic effect of morphine, and, therefore, the dose of morphine can
20 be reduced, while providing an analgesic effect equivalent to administering a higher dose of morphine alone. Such a coadministration of active agents does not effect the cataleptic properties of morphine. The reduced dose of morphine also reduces
25 adverse side effects associated with morphine administration, and can significantly reduce the addiction potential of morphine in susceptible individuals.

The present invention also is directed to
30 the administration of an endothelin antagonist to an

- 9 -

individual undergoing an opiate analgesic therapy to reduce or reverse opiate analgesic tolerance in the individual. The administration of an endothelin antagonist allows the dose of opiate analgesic to remain constant, or to be reduced, while maintaining the desired pain-reducing effect. By reducing or reversing tolerance to an opiate analgesic, the occurrence of adverse side effects can be reduced, and the possibility of opiate analgesic dependence is reduced.

The present invention, therefore, provides compositions and methods of potentiating the analgesic properties of an opiate analgesic, and of reducing or reversing tolerance to opiate analgesics. The present invention also provides pharmaceutical compositions comprising an opiate analgesic and an endothelin antagonist. Further provided are articles of manufacture comprising an opiate analgesic and an endothelin antagonist, packaged separately or together, and an insert having instructions for using the active agents.

The methods described herein benefit from the use of an opiate analgesic and an endothelin antagonist in the treatment and management of pain. The analgesic and antagonist can be administered simultaneously or sequentially to achieve the desired effect of pain treatment or reduction or reversal of opiate analgesic tolerance.

For the purposes of the invention disclosed herein, the term "treatment" includes preventing, lowering, or eliminating pain. As such,

- 10 -

the term "treatment" includes both medical therapeutic and/or prophylactic administration, as appropriate.

5 The term "cataleptic" is defined herein as a trance-like condition wherein a mammal's limbs remain in any position in which they are placed, and there is an apparent loss of sensation and awareness.

10 The term "container" means any receptacle and closure therefore suitable for storing, shipping, dispensing, and/or handling a pharmaceutical product.

15 The term "insert" means information accompanying a product that provides a description of how to administer the product, along with the safety and efficacy data required to allow the physician, pharmacist, and patient to make an informed decision regarding use of the product. The package insert generally is regarded as the "label"

20 for a pharmaceutical product.

The phrase "reducing or reversing opiate analgesic tolerance" is defined as the ability of a compound to reduce the dosage of an opiate analgesic administered to an individual to maintain a level of

25 pain control previously achieved using a greater dosage of opiate analgesic.

Several neurotransmitter mechanisms have been proposed as playing a role in the action of morphine and morphine tolerance and dependence.

30 Evidence exists that a central endothelin (ET) mechanism is involved in the actions of morphine.

- 11 -

It has been found that ET antagonists, including BQ123, for example, can potentiate morphine-induced analgesia and hyperthermia without affecting catalepsy. The present invention, therefore, provides a novel method of managing pain, reducing dependence on opioids, and reducing tolerance to opioids.

Pharmacological agents can control most pain, but it is essential to select the correct analgesic for the individual. Morphine is a major drug for the treatment of moderate to severe pain (Foley, 1993). Morphine primarily is used to treat severe pain associated with trauma, myocardial infarction, and cancer. The use of morphine in the treatment of chronic pain is limited because of inadequate analgesia, for example. Although, morphine is one of the most effective painkillers, effective pain management requires that adequate analgesia be achieved without excessive adverse side effects. Many patients treated with morphine are not successfully treated because of excessive adverse side effects and/or inadequate analgesia.

Management of excessive adverse side effects associated with morphine administration remains a major clinical challenge. Numerous strategies have been advanced to address this problem, such as (i) switching opioids, (ii) switching routes of opioid administration, (iii) improved opioid formulations, (iv) clonidine treatment, and (v) coadministering opioids that act on different receptors.

- 12 -

A massive research effort directed to the development of opioid analgesics, resulted in the discovery of numerous compounds having a varying affinity and efficacy at all the known opioid
5 receptor subtypes. Although compounds of extremely high potency have been produced, the problem of tolerance to, and dependence on, these agonists persists (Williams et al., 2001).

For example, the chronic administration of
10 morphine results in the development of physical dependence, as evidenced by the appearance of distressing physical symptoms induced by abrupt termination of morphine treatment. The signs and symptoms simulate a severe cold, and usually include
15 nasal discharge, lacrimation, chills, goose pimples, muscular aches, enhanced motor reflexes, profound body water loss attributed to hyperthermia, hyperventilation, emesis, and diarrhea (Himmelsbach, 1943; Katz et al., 1986; Maldonado et al., 1996;
20 Quock et al., 1968). It is well known that various types of opioid receptors are involved in the development of the psychological and physical dependence on opioids.

The opioid receptors have been classified
25 as μ , δ , and κ receptors, based on the relative affinity shown for experimental opioid receptor ligands. μ -Opioid receptors have been reported to play a dominant role in several pharmacological effects of morphine.

- 13 -

Role of μ -opioid receptors

An intracerebroventricular (i.c.v) injection of a selective and irreversible μ -opioid receptor antagonist, i.e., β -funaltrexamine (β -FNA),
5 drastically antagonizes morphine induced antinociception (Portoghese et al., 1980; Takemori et al., 1981; Ward et al., 1982). β -FNA also inhibits the development of physical dependence on morphine in rats (Aceto et al., 1986; DeLander et al., 1984).
10 Administration of a selective μ -opioid receptor antagonist, i.e., D-Phe-Cys-Tyr-D-Trp-Arg-The-Pen-Thr-NH₂, into the lateral cerebral ventricle 72 hours after subcutaneous implantation of two 75 mg pellets of morphine in rats induces a severe withdrawal syndrome (Maldonado et al., 1992). The
15 knockout mice with deleted μ -opioid receptors display no expression of naloxone-precipitated withdrawal symptoms including jumping and body weight loss (Matthes et al., 1996). It has been demonstrated that μ -opioid receptors consist of μ_1 and μ_2
20 subtypes in the central nervous system (CNS) (Goodman et al., 1985). The irreversible μ_1 -opioid receptor antagonist naloxonazine (Pasternak et al., 1980) is useful in investigating the function of μ_1 -
25 opioid receptors. The CXBK recombinant inbred strain of mouse is deficient in μ_1 -opioid receptors, and is much less sensitive than C57BL/6 progenitor strain to the antinociceptive and locomotor effects of morphine (Moskowitz et al., 1985). In addition,
30 the incidence of naloxone-precipitated jumping is much less in morphine-dependent C57BL/6 mice (Suzuki

- 14 -

et al., 1992a). The naloxone-induced body shakes also occurred at lower doses in C57BL/6 than in CXBK mice.

Role of δ -opioid receptors

5 Studies suggest that an interaction exists between μ - and δ -opioids. It has been found that at subantinociceptive doses, δ -opioid receptor agonists modulate antinociceptive responses to μ -opioid receptor agonists in mice (Jiang et al., 1990). Morphine acts mainly at the μ -receptor sites, but also
10 can interact with δ -opioid receptors *in vivo* and *in vitro* (Narita et al., 1993). δ -Opioid receptor antagonists do not effect morphine antinociceptive action. However, the selective blockade of δ -opioid
15 receptors by naltrindole (NTI) inhibits the development of physical dependence on morphine (Suzuki et al., 1997).

Role of κ -opioid receptors

20 Increasing evidence indicates that activation of κ -opioid receptor opposes a variety of μ -opioid receptor mediated actions throughout the brain and spinal cord (Pan, 1998). Treatment with nor-binaltorphimine (nor-BNI), a selective κ -opioid receptor antagonist, when compared to naloxone, did
25 not precipitate weight loss or other withdrawal signs in morphine-dependent mice (Cowan et al., 1988). Pretreatment with nor-BNI during chronic morphine treatment displays aggravation of the

- 15 -

weight loss precipitated by naloxone in morphine-dependent mice and rats (Suzuki et al., 1992b). These studies indicate that antagonism of endogenous κ -opioidergic system apparently elicits a potentiating effect on some morphine-withdrawal signs, including weight loss. Stimulation of endogenous κ -opioidergic system therefore should attenuate morphine withdrawal symptoms. Dynorphin A has been reported to inhibit morphine withdrawal symptoms induced by naloxone precipitation or morphine discontinuation in morphine dependent animals (Suzuki et al., 1992a). However, 3,4-dichloro-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]benzeneacetamide (i.e., U-50,488H), a selective κ -opioid receptor agonist, did not suppress the development of physical dependence on morphine in rats (Fukagawa et al., 1989). This difference has been attributed to the action of dynorphin A on all three subtypes of κ -opioid receptors, while U-50,488H acts mainly on κ_1 -opioid receptor subtype (Narita et al., 2001).

In summary, μ - and δ -opioid receptors show morphine-like withdrawal symptoms, while κ -opioid agonists do not. An opposing interaction occurs between μ/δ -opioid agonists and κ -opioid agonists.

25 Role of nonopioid receptors in morphine actions

Numerous studies, including molecular and genetic approaches, suggest a substantial role of μ -opioid receptors in the development of morphine dependence and in numerous other actions of morphine.

- 16 -

However, other systems also are involved. A role for 5-HT and cholecystokinin systems, as well as N-methyl-D-aspartate (NMDA) receptors, in opioid place conditioning has been proposed (Van Ree et al.,
5 1999).

It has been widely reported that prototypical NMDA receptor antagonists dizocilpine and ketamine, which have similar affinity for NR1/NR2A and NR1/NR2B receptors (Varney et al., 1996),
10 suppress morphine-induced place preference (Avenet et al., 1997; Tzschentke et al., 1995). Evidence is accumulating that the NR2B subunit of NMDA receptors in the nucleus accumbens may be involved in the rewarding effect of morphine (Standaert et al., 1994;
15 Watanabe et al., 1993). Neuroadaptive changes in specific brain regions that generate opioid dependence have been identified as noradrenergic transmission originating in the locus ceruleus, and most likely play the primary causal role in the expres-
20 sion of physical dependence on opioids. In contrast, a combination of behavioral and neurobiological studies point to the mesolimbic dopaminergic pathway projecting from the ventral tegmental area to the nucleus accumbens as a critical site for the
25 initiation of psychological dependence on opioids.

Endothelin in the central nervous system

ET is an extremely potent endothelium derived vasoconstriction factor (Hickey et al., 1985) that was isolated, sequenced, and cloned (Yanagisawa
30 et al., 1988). Endothelins are 21 amino acid, high-

- 17 -

ly potent vasoconstrictive peptides with two disulfide bonds. Endothelins are produced biologically by enzymatically cleaving preproendothelin to proendothelin, then to endothelin by endothelin-
5 converting enzymes. ET exerts biological effects by binding to cell surface receptors which are 7-transmembrane receptors coupled to G-proteins. There are two distinct types of endothelin receptors, (a) the ET-1 selective ET_A receptors primarily found on
10 vascular smooth muscle and responsible for vasoconstriction, and (b) nonselective ET_B receptors primarily found in vascular endothelium and responsible for vasodilation.

The vasoconstrictive effects of ET-1 are
15 mediated predominantly by G-protein coupled ET_A receptors (Reynolds et al., 1989). ET-1 also is made in high concentrations by prostate, metastatic cancers, and CNS. ET in the CNS is produced by endothelial cells and nonendothelial cells, such as
20 neurons, astrocytes, and glial cells (MacCumber et al., 1990).

The global distribution of ET and its binding sites in the brain suggests that, in addition to being a vasoconstrictor, it may be acting as
25 an important neuropeptide in the CNS (Gulati et al., 1992). Endothelin (ET) receptor antagonists, in particular selective ET_A or balanced antagonists ET_A/ET_B, represent a therapeutic area for diseases such as congestive heart failure (CHF) and pulmonary
30 hypertension. BQ-123 and BMS-182874 are specific antagonists of ET_A receptors (Ihara et al., 1992;

- 18 -

Stein et al., 1994). Endothelin antagonists have profound effects on the pulmonary vasculature and the right heart, whereas ACE inhibitors primarily affect the peripheral vessel and the left heart.

5 Several studies indicate that the central ET receptors are predominantly of ET_B subtype (Matsumura et al., 1991). Rat cerebral astrocytes have been shown to express mainly ET_B type of receptors (Hama et al., 1992) and glial cells also
10 were found to intensely express ET_B receptor mRNA (Pagotto et al., 1995). However, the central administration of a highly selective ET_B receptor agonist, IRL-1620, does not produce any effect on the cardiovascular system, and the systemic and
15 regional circulatory effects of centrally administered ET-1 have been shown to be mediated through the ET_A receptors (Gulati et al., 1995; Rebello et al., 1995).

 Intracerebroventricular administration of
20 ET-1 produces a transient rise followed by sustained fall in the mean arterial blood pressure (BP) (Gulati et al., 1996). The pressor effect was accompanied by an increase in renal sympathetic nerve activity and plasma levels of catecholamines
25 and arginine-vasopressin (Matsumura et al., 1991).

 It also has been shown that the effects of central administration of ET-1 are mediated through activation of the sympathetic nervous system because these effects were attenuated by ganglion blockers
30 (Kawano et al., 1989; Matsumura et al., 1991). Intracisternal administration of ET-1 elicited a

- 19 -

transient increase in BP, renal sympathetic nerve activity, and phrenic nerve activity. A subsequent fall in BP was accompanied by a decrease in renal sympathetic nerve activity and phrenic nerve activity (Kuwaki et al., 1994). The observation that central ET-1 induced increase in pressor response was suppressed by pretreatment with phenoxybenzamine (Ouchi et al., 1989), further implicates the active participation of sympathetic nervous system in the initial pressor phase.

Interaction of opioids and endothelin

Both ET and opioids exist in the CNS and can play a role in regulation of cardiovascular and/or other functions. These two peptides can have some interactions, especially through the sympathetic nervous system because both are located in sites involved in cardiovascular regulation, such as, for example, the hypothalamus, nucleus tractus solitarius, and locus ceruleus. Very few studies exist indicating that ET and opioids interact and modulate the effects of one another.

For example, it has been reported that ET can induce pain, which can be blocked by morphine. ET-1 administered intraperitoneally into Cr1:CD-1(ICR)BR mice and CXBK mice produced abdominal irritation which produced a single occurrence of a wave of constriction and elongation passing caudally along the abdominal wall accompanied by a mild twisting of the trunk followed by extension of the hind limbs. This ET-1 induced nociception could be

- 20 -

blocked by morphine through μ_1 sensitive pathway
(Raffa et al., 1994). In another study, ET-1 (200
to 800 μ M) applied to rat sciatic nerve produced
reliable, robust, unilateral hindpaw flinching
5 lasting for 60 min. Pretreatment with morphine
completely blocked this effect in a naloxone
sensitive manner. BQ123, an ET_A receptor antag-
onist, also blocked the ET-1 induced hind paw
flinching (Davar et al., 1998). The role of ET_A
10 receptor antagonist ABT-627 in tactile allodynia has
been investigated in streptozotocin-induced diabetic
rat model of neuropathic pain.

The systemic administration of ABT-627
produced a dose-dependent increase (40 to 50%) in
15 tactile allodynia thresholds. The antinociceptive
effect of ABT-627 was maintained following chronic
administration of the antagonist in drinking water
for 7 days. In comparison, morphine produced a
significant (90%) increase in tactile allodynia
20 thresholds. ET_B receptor antagonist did not affect
the tactile allodynia threshold (Jarvis et al.,
2000).

Responses to morphine can be associated
with increased systemic and cerebrovascular levels
25 of ET-1 and upregulation of ET-1 and ET_A receptor
mRNA in the brainstem of newborn piglets (Modanlou
et al., 1998). ET-1 administered intracerebroven-
tricularly (i.c.v.) caused significant increases in
mean arterial pressure and RSNA, and these effects
30 were potentiated by naloxone pretreatment (Matsumura
et al., 1994). The effect is centrally mediated

- 21 -

because naloxone methobromide, a naloxone derivative that does not cross the blood-brain barrier, did not alter the baroreflex sensitivity. It was concluded that ET-1 exerts a potent central pressor action
5 mediated by enhanced sympathoadrenal outflow, and naloxone potentiates these pressor and sympathetic responses.

In accordance with an important feature of the present invention, it has been hypothesized that
10 actions of morphine and related opioids mediated through the sympathetic pathway can be modulated by ET antagonists, and that an interaction exists between opioids and ET. This approach is particularly useful for the management of symptoms of
15 morphine withdrawal.

In accordance with another important feature of the present invention, an opiate analgesic is present in a composition, or is administered, with an endothelin antagonist in a weight ratio of
20 analgesic-to-antagonist of about 0.01:1 to about 100:1, preferably about 0.02:1 to about 50:1, and most preferably about 0.1:1 to about 10:1. This ratio depends upon the type and identity of opioid analgesic and endothelin antagonist being used. The
25 ratio of analgesic-to-antagonist that is administered is dependent upon the particular analgesic and antagonist used, and the origin and severity of the pain being treated. This ratio can be readily
30 determined by a person skilled in the art to achieve the desired reduction in pain.

- 22 -

An opiate analgesic utilized in the present invention can be one or more opium alkaloid or semisynthetic opiate analgesic. Specific opiate analgesics include, but are not limited to, (a) opium; (b) opium alkaloids, such as morphine, morphine sulfate, codeine, codeine phosphate, codeine sulfate, diacetylmorphine, morphine hydrochloride, morphine tartrate, and diacetylmorphine hydrochloride; and (c) semisynthetic opiate analgesics, such as dextromethorphan hydrobromide, hydrocodone bitartrate, hydromorphone, hydromorphone hydrochloride, levorphanol tartrate, oxymorphone hydrochloride, and oxycodone hydrochloride. Other opioids include, but are not limited to, fentanyl, meperidine, methodone, and propoxyphene.

An endothelin antagonist utilized in the present invention can be any of the endothelin receptor antagonists known in the art. Endothelin is a potent vasoconstrictor. Endothelin antagonists are used to treat acute heart failure, congestive/chronic heart failure, pulmonary arterial hypertension, pulmonary edema, subarachnoid hemorrhage, chronic obstructive pulmonary disease, myocardial infarction, acute cerebral ischemia, acute coronary syndromes, acute renal failure, post-operative treatment in liver operations, and prostate cancer.

No adverse effects are expected when a healthy patient is administered an opiate analgesic in combination with an endothelin antagonist. However, for patients suffering from conditions like congestive heart failure and other diseases treat-

- 23 -

able by an endothelin antagonist, coadministration of an opioid analgesic and an endothelin antagonist should be monitored carefully.

Preferred ET antagonists are antagonists
5 selective for endothelin A (ET_A) receptors or are
balanced ET_A/endothelin B (ET_B) antagonists. Such ET
antagonists are set forth in Appendices A and B
herein. However, endothelin B antagonists and
miscellaneous endothelin antagonists, as set forth
10 in Appendices C and D herein, also can be used in a
composition or method of the present invention.
Additional useful endothelin antagonists can be
found in U.S. Patent Application Publication No.
US 2002/0082285 A1, incorporated herein by refer-
15 ence.

Specific examples of endothelin antag-
onists useful in the present invention include, but
are not limited to, atrasentan, tezosentan, bosen-
tan, sitaxsentan, enrasentan, BMS-207940 (Bristol-
20 Myers Squibb), BMS-193884, BMS-182874, J-104132
(Banyu Pharmaceutical), VML 588/Ro 61-1790 (Vanguard
Medica), T-0115 (Tanabe Seiyaku), TAK-044 (Takeda),
BQ-788, BQ123, YM-598, LU 135252, PD 145065,
A-127722, ABT-627, A-192621, A-182086, TBC3711,
25 BSF208075, S-0139, TBC2576, TBC3214, PD156707,
PD180988, ABT-546, ABT-627, Z1611, RPR118031A,
SB247083, SB217242, S-Lu302872, TPC10950, and
SB209670.

BQ123 is a specific endothelin A antag-
30 onist, and is the sodium salt of cyclo(-D-Trp-D-Asp-
Pro-D-Val-Leu-). BQ-788 is a specific endothelin B

- 24 -

antagonist, and is the sodium salt of N-cis-2,6-dimethylpiperidinocarbonyl-L-gamma-methyllleucyl-D-1-methoxycarbonyl triptophanyl-DNIE (see *Proc. Natl. Acad. Sci. USA*, 91, pp. 4892-4896 (1994)).

5 In addition to a conventional endothelin antagonist, a compound that inhibits the formation of endogenous endothelin also can be used as the endothelin antagonist in the present invention. Such compounds are useful because they prevent endothelin formation, and, therefore, decrease the
10 activity of endothelin receptors. One class of such compounds is the endothelin converting enzyme (ECE) inhibitors.

 Useful ECE inhibitors include, but are not
15 limited to, CGS34225 (i.e., N-((1-((2(S)-(acetylthio)-1-oxopentyl)-amino)-1-cyclopentyl)-carbonyl-S-4-phenylphenyl-alanine methyl ester) and phosphoramidon (i.e., N-(α -rhamnopyranosyloxyhydroxyphosphinyl)-Leu-Trp).

20 The following tests were conducted to illustrate the potentiating effects of an endothelin antagonist on an opiate analgesic administered to a mammal, including humans.

 Male Sprague-Dawley rats weighing 225 to
25 250 g (Sasco King Animal Co., Madison, WI) were housed in a room with controlled temperature (23 \pm 1°C), humidity (50 \pm 10%), and light (6:00 a.m. to 6:00 p.m.) for at least four days prior to testing. Food and water were available to the rats continuously.
30

- 25 -

The rats were divided into four groups (Figure 1): group 1 received vehicle (saline 5 μ l, i.c.v. over 5 minutes) and vehicle (saline 100 μ l/kg s.c.); group 2 received vehicle (saline 100 μ l/kg, s.c.) and BQ123 (10 μ g, i.c.v. in a volume of 5 μ l over 5 minutes); group 3 received morphine (8 mg/kg, s.c. in a volume of 100 μ l/kg) and vehicle (saline 100 μ l/kg, s.c.); and group 4 received morphine (8 mg/kg, s.c. in a volume of 100 μ l/kg) and BQ123 (10 μ g, i.c.v. in a volume of 5 μ l over 5 minutes). Vehicle or BQ123 treatment was performed 30 minutes prior to morphine administration.

Tests also were performed on mice. The mice were housed and fed similar to the rats as described above taking into consideration the size differential between rats and mice. The mice were divided into groups identical to the rats, but administered different amounts of vehicle, opiate analgesic, and endothelin antagonist, as disclosed hereafter.

Measurement of analgesic effect: The analgesic response to morphine was determined by the tail flick method. The tail flick latencies to thermal stimulation were determined before and 30, 60, 90, 120, 180, 210, 240, 270, 300, and 360 minutes after the morphine injection. The basal tail flick latency was about 2 seconds. A cutoff time of 10 seconds was used to prevent damage to the tail. The basal latency was subtracted from that induced by morphine. The analgesic response in each rat was converted into $AUC_{0 \rightarrow 360 \text{ min}}$. (area under the

- 26 -

curve), and was expressed as mean \pm SEM. Eight to nine animals were used for each dose of morphine sulfate. The differences in analgesic response to different doses of morphine were compared by using student's t test. A value of $P < 0.05$ was considered significant.

Measurement of colonic temperature: The change in temperature in response to morphine was determined. The colonic temperature of each rat was recorded before and various times after morphine injection for a period of 360 minutes using a telethermometer. The change in temperature from the basal value was plotted with time and was converted to $AUC_{0 \rightarrow 300 \text{ min.}}$ and was expressed as mean \pm SEM. Eight to nine rats were used for each dose of morphine sulfate. Rats used to measure tail flick latencies also were used to determine the colonic temperature. The differences in temperature response between various groups were compared by using student's t test. A value of $P < 0.05$ was considered significant.

Measurement of catalepsy: Rats were tested for catalepsy by the bar test. An aluminum bar 5 mm in diameter was placed 4 cm above the floor, and the animal's forepaw was gently placed on the bar. The time required for the animal to place at least one paw on the floor was measured with a maximum term of observation of 180 seconds. Behavioral evaluation of catalepsy was performed 45 minutes after the administration of morphine or its vehicle. Statistical tests were performed as described above.

- 27 -

Effect on body weight: Body weight was measured in all groups of rats of Fig. 1. BQ123 pretreatment induced no change in body weight between the four test groups. Body weight was
5 measured before administering any drug and at the end of the experiment, i.e., seven hours after the administration of morphine. The data clearly indicates that all groups were very comparable in their response to change in body weight, and that
10 neither morphine nor BQ123 produced any acute effect on body weight.

Effect on cataleptic behavior: Morphine administration significantly increased catalepsy in rats. In contrast, BQ123 did not produce any cata-
15 leptic effect. Catalepsy is an overall indication of motor behavior including locomotor activity of rats. Pretreatment with BQ123 did not produce any change in morphine-induced catalepsy. In particular, morphine produced a significant increase in
20 catalepsy compared to control group 1, but this increase in catalepsy was not affected by BQ123 pretreatment. Therefore, the effect of morphine on cataleptic behavior was not affected by administration of the endothelin antagonist BQ123. These
25 findings indicate that BQ123 did not affect a pharmacological action produced in rats by morphine.

Effect on body temperature: The control treated rats (group 1 of Fig. 1) did not show any effect on body temperature. BQ123 treatment also
30 did not produce any significant effect on body temperature. Morphine treatment produced signifi-

- 28 -

cant hyperthermia in rats. Hyperthermia lasted for about four hours after the administration of morphine. The hyperthermic effect of morphine was significantly potentiated by BQ123. The hyperthermic effect of morphine was not only significantly greater in BQ123 pretreated rats, but lasted for more than six hours. In particular, it was demonstrated that morphine produced a significant increase in body temperature compared to control group 1. Pretreatment with BQ123 significantly potentiated the effect of morphine-induced hyperthermia compared to vehicle and morphine-treated rats. This is a significant observation because although morphine-induced catalepsy is not affected by endothelin antagonist BQ123, the hyperthermic response to morphine is potentiated by BQ123.

Effect on analgesia: The control group of rats exhibited tail flick latencies of about 2 seconds. BQ123 treatment did not produce any significant effect on tail flick latency. Morphine (8 mg/kg, s.c.) produced significant analgesia in rats, and the tail flick latencies reached more than 10 seconds (Figure 2). A significant increase in tail flick latencies was observed until three hours after the administration of morphine. The analgesic effect of morphine was significantly potentiated by BQ123. The analgesic effect of morphine not only was significantly greater in BQ123 pretreated rats, but lasted for more than six hours.

The effect of BQ123 treatment (3 μ g, i.c.v.) on morphine analgesia also was determined in

- 29 -

mice (Figures 3-5). Morphine (2, 4, and 8 mg/kg, s.c.) produced a significant increase in mouse tail flick latency that lasted for 1.5 hours. BQ123 significantly potentiated morphine analgesia, which
5 lasted for more than 4 hours. Figure 3 especially shows that tail flick latency is increased by administration of BQ123, thereby showing an endothelin antagonist potentiating effect on morphine at a low dosage administration of the opiate analgesic.

10 Figures 6-8 illustrate the ability of an endothelin antagonist to induce opiate analgesic tolerance in mice and rats rendered tolerant and dependent on morphine. The test method used to generate the data set forth in Figs. 6-8 is disclosed in H.N. Bhargava et al., *J. Pharmacol. Exp. Ther.*, 252(3), pp. 901-907 (1990). The method disclosed therein was modified slightly for mice because the smaller mice could not survive the morphine dose administered to the rats. Tolerance to
15 morphine is demonstrated in Figs. 6-8 for the endothelin antagonists BQ123 and BMS182874.

The effect of BQ123 (3 µg, i.c.v.) on naloxone-precipitated morphine withdrawal also was determined in mice. Naloxone (1 mg/kg, i.p.) administration to morphine tolerant mice (1 morphine 75 mg pellet for 3 days) resulted in expression of withdrawal symptoms. BQ123 did not affect hypothermia, loss of body weight, jumping behavior, falls, diarrhea, fecal boli, urination, ptosis,
25 writhing, and rearing behavior during withdrawal. Tests also were performed to determine tolerance to

- 30 -

analgesic effect in morphine tolerant dependent rats (6 morphine pellets in 7-day period). BQ123-treated (10 µg, i.c.v., twice a day for 7 days) rats did not become tolerant to morphine.

5 These studies demonstrate that an endothelin antagonist, like BQ123, potentiates the analgesic effect of morphine and prevents the development of tolerance to morphine analgesia, without affecting naloxone-precipitated morphine withdrawal.
10 The combined use of BQ123 or other ET antagonist and an opiate provides a novel approach to improving analgesia and eliminating morphine tolerance. These findings provide a novel combination of active agents to manage various types of pain.

15 In summary, rat and mice studies showed that endothelin antagonist administration in combination with an opiate analgesic potentiated analgesia and hypothermia, but did not affect catalepsy or body weight. A mouse study administering BQ123
20 and morphine (2, 4, or 8 mg/kg) also showed that analgesia was potentiated, but hypothermia and body weight were not affected.

 Another study directed to naloxone (1 mg/kg) initiated withdrawal symptoms in mice in
25 conjunction with a combination morphine and BQ123 treatment showed no effect on temperature, body weight, urination, diarrhea, jumping behavior, number of falls, ptosis, and grooming.

 Overall, test results show that an endothelin antagonist significantly potentiates morphine
30 analgesia in mice and rats, prevents the development

- 31 -

of tolerance to analgesic actions of opiate anal-
gesics, like morphine, and potentiates morphine-
induced hyperthermia in rats, but does not affect
morphine-induced hypothermia in mice, does not
5. affect morphine-induced catalepsy in rats, and does
not affect naloxone precipitated morphine with-
drawal. Therefore, an endothelin antagonist poten-
tiates morphine analgesia, but does not potentiate
morphine withdrawal symptoms and prevents or re-
10 verses the development of tolerance to analgesic
opiates.

The test results clearly demonstrate that
endothelin antagonists, like BQ123, potentiate mor-
phine-induced analgesia and hyperthermia without
15 affecting morphine-induced cataleptic behavior.
This is an important clinical finding because endo-
thelin antagonists have minimal cardiovascular
effects in normal healthy individuals and because ET
antagonists, like tezosentan, bosentan, darnsentan,
20 and atrasentan, are nearing regulatory approval.
Endothelin antagonists combined with morphine,
therefore, can be used to potentiate the analgesic
action of morphine without affecting some of the
other pharmacological actions of morphine.

25 It has been demonstrated that ET signif-
icantly regulates the central autonomic nervous
system (Gulati et al., 1997; Kumar et al., 1997),
and most of the withdrawal reactions of morphine
also are mediated through the central autonomic
30 nervous system. Central ET modulates pharmacolog-
ical actions of morphine. Using an ET antagonist

- 32 -

together with morphine increases the analgesic and hyperthermic action of morphine, but does not affect the cataleptic action of morphine. On the basis of results obtained, an ET antagonist can reduce the
5 dose of morphine and still produce same degree of analgesic action of morphine as produced by a higher dose of morphine used alone. Lowering the dose of morphine can significantly reduce the addiction potential of morphine in patients.

10 These findings show that when combined with an endothelin antagonist, morphine and other opiate analgesics produce significant analgesia with a much lower dose of analgesic, and, therefore, the addiction potential of the opiate analgesic is re-
15 duced. These observations also indicate that the duration of analgesic response of morphine can be significantly increased by administration of an endothelin antagonist.

20 The data shows that some morphine-induced pharmacological responses, like change in temperature and analgesia, can be separately potentiated while other responses, like catalepsy, are not affected by endothelin antagonist administration.

25 The above tests and data show that a combination of an opiate analgesic and an endothelin antagonist can be administered to mammals in methods of treating pain. The opiate analgesic and endothelin antagonist can be formulated in suitable excipients for oral administration, or for parenteral
30 al administration. Such excipients are well known in the art. The active agents typically are present

- 33 -

in such a composition in an amount of about 0.1% to about 75% by weight, either alone or in combination.

Pharmaceutical compositions containing the active agents, i.e., opiate analgesic and endothelin antagonist, of the present invention are suitable for administration to humans or other mammals. Typically, the pharmaceutical compositions are sterile, and contain no toxic, carcinogenic, or mutagenic compounds that would cause an adverse reaction when administered.

The method of the invention can be accomplished using the active agents as described above, or as a physiologically acceptable salt or solvate thereof. The active agents, salts, or solvates can be administered as the neat compounds, or as a pharmaceutical composition containing either or both entities.

The active agents can be administered by any suitable route, for example by oral, buccal, inhalation, sublingual, rectal, vaginal, intracisternal through lumbar puncture, transurethral, nasal, percutaneous, i.e., transdermal, or parenteral (including intravenous, intramuscular, subcutaneous, and intracoronary) administration. Parenteral administration can be accomplished using a needle and syringe, or using a high pressure technique, like POWDERJECT™. Administration of the active agents can be performed before, during, or after the onset of pain.

The pharmaceutical compositions include those wherein the active ingredients are adminis-

- 34 -

tered in an effective amount to achieve their intended purpose. More specifically, a "therapeutically effective amount" means an amount effective to prevent development of, to eliminate, or to
5 alleviate pain. Determination of a therapeutically effective amount is well within the capability of those skilled in the art, especially in light of the detailed disclosure provided herein.

A "therapeutically effective dose" refers
10 to the amount of the active agents that results in achieving the desired effect. Toxicity and therapeutic efficacy of such active agents can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g.,
15 determining the LD₅₀ (the dose lethal to 50% of the population) and the ED₅₀ (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index, which is expressed as the ratio be-
20 tween LD₅₀ and ED₅₀. A high therapeutic index is preferred. The data obtained from such data can be used in formulating a range of dosage for use in humans. The dosage of the active agents preferably lies within a range of circulating concentrations
25 that include the ED₅₀ with little or no toxicity. The dosage can vary within this range depending upon the dosage form employed, and the route of administration utilized.

The exact formulation, route of adminis-
30 tration, and dosage is determined by an individual physician in view of the patient's condition. Dos-

- 35 -

age amounts and intervals can be adjusted individually to provide levels of active agents that are sufficient to maintain therapeutic or prophylactic effects.

5 The amount of active agents administered is dependent on the subject being treated, on the subject's weight, the severity of the affliction, the manner of administration, and the judgment of the prescribing physician.

10 Specifically, for administration to a human in the curative or prophylactic treatment of pain, oral dosages of an opiate analgesic and endothelin antagonist, individually generally are about 10 to about 200 mg daily for an average adult
15 patient (70 kg), typically divided into two to three doses per day. Thus, for a typical adult patient, individual tablets or capsules contain about 0.1 to about 200 mg opioid analgesic and about 0.1 to about 50 mg endothelin antagonist, in a suitable pharmaceutically acceptable vehicle or carrier, for administration in single or multiple doses, once or
20 several times per day. Dosages for intravenous, buccal, or sublingual administration typically are about 0.1 to about 10 mg/kg per single dose as required. In practice, the physician determines the actual dosing regimen that is most suitable for an individual patient, and the dosage varies with the age, weight, and response of the particular patient. The above dosages are exemplary of the average case,
25 but there can be individual instances in which
30

- 36 -

higher or lower dosages are merited, and such are within the scope of this invention.

The active agents of the present invention can be administered alone, or in admixture with a pharmaceutical carrier selected with regard to the
5 intended route of administration and standard pharmaceutical practice. Pharmaceutical compositions for use in accordance with the present invention thus can be formulated in a conventional manner using one
10 or more physiologically acceptable carriers comprising excipients and auxiliaries that facilitate processing of the active agents into preparations that can be used pharmaceutically.

These pharmaceutical compositions can be
15 manufactured in a conventional manner, e.g., by conventional mixing, dissolving, granulating, dragee-making, emulsifying, encapsulating, entrapping, or lyophilizing processes. Proper formulation is dependent upon the route of administration
20 chosen. When a therapeutically effective amount of the active agents are administered orally, the composition typically is in the form of a tablet, capsule, powder, solution, or elixir. When administered in tablet form, the composition can additionally
25 contain a solid carrier, such as a gelatin or an adjuvant. The tablet, capsule, and powder contain about 5% to about 95% of an active agent of the present invention, and preferably from about 25% to about 90% of an active agent of the present in-
30 vention. When administered in liquid form, a liquid carrier, such as water, petroleum, or oils of animal

- 37 -

or plant origin, can be added. The liquid form of the composition can further contain physiological saline solution, dextrose or other saccharide solutions, or glycols. When administered in liquid
5 form, the composition contains about 0.5% to about 90% by weight of active agents, and preferably about 1% to about 50% of an active agents.

When a therapeutically effective amount of the active agents is administered by intravenous,
10 cutaneous, or subcutaneous injection, the composition is in the form of a pyrogen-free, parenterally acceptable aqueous solution. The preparation of such parenterally acceptable solutions, having due regard to pH, isotonicity, stability, and the like,
15 is within the skill in the art. A preferred composition for intravenous, cutaneous, or subcutaneous injection typically contains, in addition to a compound of the present invention, an isotonic vehicle.

Suitable active agents can be readily
20 combined with pharmaceutically acceptable carriers well-known in the art. Such carriers enable the active agents to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a
25 patient to be treated. Pharmaceutical preparations for oral use can be obtained by adding the active agents with a solid excipient, optionally grinding the resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if
30 desired, to obtain tablets or dragee cores. Suitable excipients include, for example, fillers and

- 38 -

cellulose preparations. If desired, disintegrating agents can be added.

The active agents can be formulated for parenteral administration by injection, e.g., by
5 bolus injection or continuous infusion. Formulations for injection can be presented in unit dosage form, e.g., in ampules or in multidose containers, with an added preservative. The compositions can take such forms as suspensions, solutions, or emul-
10 sions in oily or aqueous vehicles, and can contain formulatory agents, such as suspending, stabilizing, and/or dispersing agents.

Pharmaceutical compositions for parenteral administration include aqueous solutions of the
15 active agent in water-soluble form. Additionally, suspensions of the active agents can be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils or synthetic fatty acid esters. Aqueous injection
20 suspensions can contain substances which increase the viscosity of the suspension. Optionally, the suspension also can contain suitable stabilizers or agents that increase the solubility of the compounds and allow for the preparation of highly concentrated
25 solutions. Alternatively, a present composition can be in powder form for constitution with a suitable vehicle, e.g., sterile pyrogen-free water, before use.

The active agents also can be formulated
30 in rectal compositions, such as suppositories or retention enemas, e.g., containing conventional

- 39 -

suppository bases. In addition to the formulations described previously, the active agents also can be formulated as a depot preparation. Such long-acting formulations can be administered by implantation
5 (for example, subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, the active agents can be formulated with suitable polymeric or hydrophobic materials (for example, as an emulsion in an acceptable oil) or ion exchange
10 resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

In particular, the active agents can be administered orally, buccally, or sublingually in the form of tablets containing excipients, such as
15 starch or lactose, or in capsules or ovules, either alone or in admixture with excipients, or in the form of elixirs or suspensions containing flavoring or coloring agents. Such liquid preparations can be prepared with pharmaceutically acceptable additives,
20 such as suspending agents. An active agent also can be injected parenterally, for example, intravenously, intramuscularly, subcutaneously, intrathecally, intracisternally, or intracoronarily. For parenteral administration, the active agent is best used
25 in the form of a sterile aqueous solution which can contain other substances, for example, salts, or monosaccharides, such as mannitol or glucose, to make the solution isotonic with blood.

For veterinary use, the active agents are
30 administered as a suitably acceptable formulation in accordance with normal veterinary practice. The

- 40 -

veterinarian can readily determine the dosing regimen and route of administration that is most appropriate for a particular animal.

As stated above, morphine is one of the most potent analgesics, and is widely used for pain management in several disease conditions, including cancer. A major problem in the use of morphine, and other opiate analgesics, is their potential to produce sedation tolerance/dependent, and respiratory depressions, and to cause addiction.

It has been discovered that using an endothelin antagonist in combination with an opiate analgesic potentiates the analgesic and hyperthermic action of the analgesic, but does not increase the sedative or cataleptic action of the analgesic. The combined opiate analgesic-endothelin antagonist treatment can be used, for example, in office surgeries, oral surgeries, and post-surgical pain management.

It also has been discovered that by using an endothelin antagonist together with an opiate analgesic, the dose of morphine can be reduced, and still provide the same analgesic action as a larger dose of morphine used alone. By using less morphine, the addiction potential of an opiate analgesic in patients can be reduced significantly. The administration of an endothelin antagonist to an individual undergoing opiate analgesic treatment, therefore, reduces or eliminates tolerance to opiate analgesics.

- 41 -

References:

- M.D. Aceto et al., *Eur J Pharmacol*,
123(3):387-93 (1986).
- P. Avenet et al., *Neurosci Lett*,
5 223(2):133-6 (1997).
- A. Cowan et al., *J Pharmacol Exp Ther*,
246(3):950-5 (1988).
- G. Davar et al., *Neuroreport*, 9(10):2279-
83 (1998).
- 10 G.E. DeLander et al., *J Pharmacol Exp
Ther*, 231(1):91-6 (1984).
- K.M. Foley, *Handbook of Experimental
Pharmacology*, Springer-Verlag, Berlin (1993).
- Y. Fukagawa et al., *Eur J Pharmacol*,
15 170(1-2):47-51 (1989).
- R.R. Goodman et al., *Proc Natl Acad Sci
USA*, 82(19):6667-71 (1985).
- F.J. Gordon et al., *J Pharmacol Exp Ther*,
237(2):428-36 (1986).
- 20 F.J. Gordon, *Peptides*, 11(2):305-9 (1990).
- A. Gulati et al., *Life Sci*, 58(5):437-45
(1996).
- A. Gulati et al., *Life Sci*, 51(22):1715-24
(1992).
- 25 A. Gulati et al., *Am J Physiol*, 273(3 Pt
2):H1177-86 (1997).
- A. Gulati et al., *J Cardiovasc Pharmacol*,
26(Suppl 3):S244-6 (1995).
- A. Gulati et al., *Drug Develop Res*,
30 26:361-387 (1992).

- 42 -

- H. Hama et al., *Biochem Biophys Res Commun*, 186(1):355-62 (1992).
- K.A. Hickey et al., *Am J Physiol*, 248(5 Pt 1):C550-6 (1985).
- 5 Himmelsbach, *Fed Proc*, 2:201-203 (1943).
- M. Ihara et al., *Life Sci*, 50(4):247-55 (1992).
- M.F. Jarvis et al., *Eur J Pharmacol*, 388(1):29-35 (2000).
- 10 Q. Jiang et al., *Prog Clin Biol Res*, 328:449-52 (1990).
- J.L. Katz et al., *Behavioral Analysis of Drug Dependence*, ed. S.R. Goldberg et al., Vol. Ch. 8, pp. 287-327, Academic Press, Orlando (1986).
- 15 Y. Kawano et al., *J Hypertens Suppl* 7(6):S22-3 (1989).
- S. Koyama et al., *Eur J Pharmacol*, 90(4):367-76 (1983).
- A. Kumar et al., *Peptides*, 18(6):855-64
- 20 (1997).
- T. Kuwaki et al., *Jpn J Physiol*, 44(1):1-18 (1994).
- M.W. MacCumber et al., *Proc Natl Acad Sci USA*, 87(6):2359-63 (1990).
- 25 R. Maldonado et al., *J Pharmacol Exp Ther*, 261(2):669-77 (1992).
- R. Maldonado et al., *Neurobiological Mechanisms of Opiate Withdrawal*, ed. R. Maldonado et al., Vol. Ch. 5, pp. 77-124, Springer, New York
- 30 (1996).

- 43 -

K. Matsumura et al., *Am J Physiol*, 266(4 Pt 2):R1403-10 (1994).

K. Matsumura et al., *Hypertension*, 19(6 Pt 2):648-52 (1992).

5 K. Matsumura et al., *Hypertension*, 17(6 Pt 2):1192-6 (1991).

H.W. Matthes et al., *Nature*, 383(6603): 819-23 (1996).

C.N. May et al., *Clin Sci (Colch)*,
10 76(4):431-7 (1989).

H.D. Modanlou et al., *Can J Physiol Pharmacol*, 76(4):443-50 (1998).

A.S. Moskowitz et al., *Brain Res*, 360(1-2):108-16 (1985).

15 M. Narita et al., *Pharmacol Ther*, 89(1):1-15 (2001).

M. Narita et al., *Psychopharmacology*, 111(4):423-6 (1993).

Y. Ouchi et al., *Am J Physiol*, 256(6 Pt 20 2):H1747-51 (1989).

U. Pagotto et al., *J Cardiovasc Pharmacol*, 26(Suppl 3):S104-6 (1995).

Z.Z. Pan, *Trends Pharmacol Sci*, 19(3):94-8 (1998).

25 G.W. Pasternak et al., *J Med Chem*, 23(6):674-6 (1980).

A. Pfeiffer et al., *Endocrinology*, 113(3):929-38 (1983).

P.S. Portoghese et al., *J Med Chem*,
30 23(3):233-4 (1980).

- 44 -

C.P. Quock et al., *Br J Addict Alcohol Other Drugs*, 63:261-270 (1968).

R.B. Raffa et al., *Life Sci*, 54(4):157-62 (1994).

5 S. Rebello et al., *Brain Res*, 676(1):141-50 (1995).

E.E. Reynolds et al., *Biochem Biophys Res Commun*, 160(2):868-73 (1989).

10 T. Sakurai et al., *Nature*, 348(6303):732-5 (1990).

D.G. Standaert et al., *J Comp Neurol*, 343(1):1-16 (1994).

P.D. Stein et al., *J Med Chem*, 37(3):329-31 (1994).

15 T. Suzuki et al., *Life Sci*, 50(12):849-56 (1992a).

T. Suzuki et al., *Eur J Pharmacol*, 213(1):91-7 (1992b).

20 T. Suzuki et al., *Pharmacol Biochem Behav*, 57(1-2):293-9 (1997).

A.E. Takemori et al., *Eur J Pharmacol*, 70(4):445-51 (1981).

T.M. Tzschentke et al., *Neurosci Lett*, 193(1):37-40 (1995).

25 J.M. Van Ree et al., *Pharmacol Rev*, 51:342-396 (1999).

M.A. Varney et al., *J Pharmacol Exp Ther*, 279(1):367-78 (1996).

30 S.J. Ward et al., *J Pharmacol Exp Ther*, 220:494-498 (1982).

- 45 -

M. Watanabe et al., *J Comp Neurol*,
338:377-390 (1993).

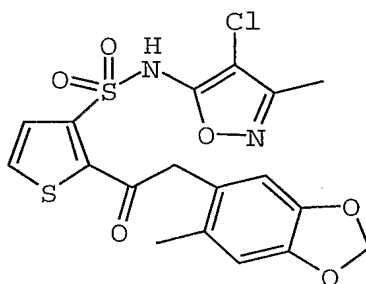
J.T. Williams et al., *Physiol Rev*,
81(1):299-343 (2001).

5 M. Yanagisawa et al., *J Hypertens Suppl*
6(4):S188-91 (1988).

Modifications and variations of the
invention as hereinbefore set forth can be made
without departing from the spirit and scope thereof,
10 and, therefore, only such limitations should be
imposed as are indicated by the appended claims.

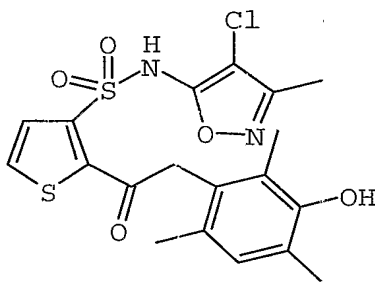
- 46 -

APPENDIX A
SELECTIVE ET_A ANTAGONISTS



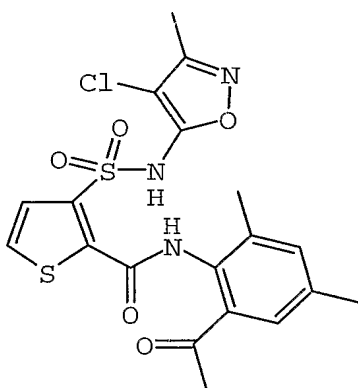
1

sitaxsentan



2

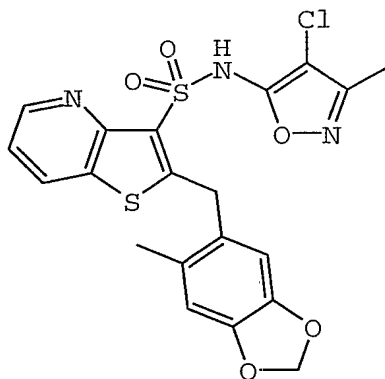
TBC2576



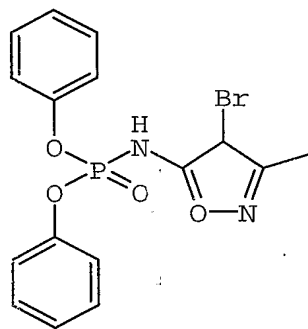
3

TBC3214

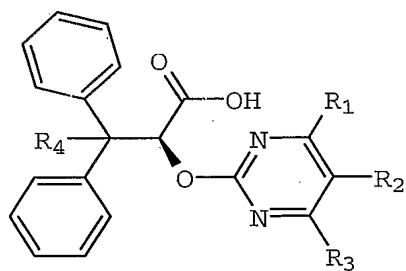
- 47 -



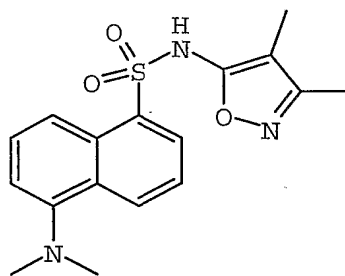
4



5

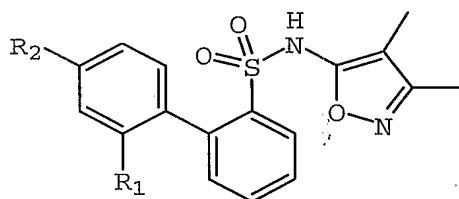
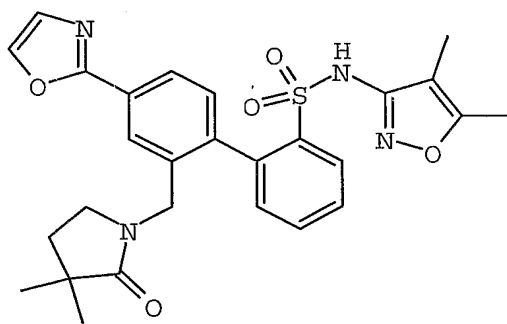
6 $R_1=R_3=R_4=CH_3$, $R_2=H$ 7 $R_1=R_3=R_4=OCH_3$, $R_2=F$ 8 $R_1=OCH_3$, $R_2=H$, $R_3=CH_3$, $R_4=-OCH_2CON(CH_3)C_6H_5$

- 48 -



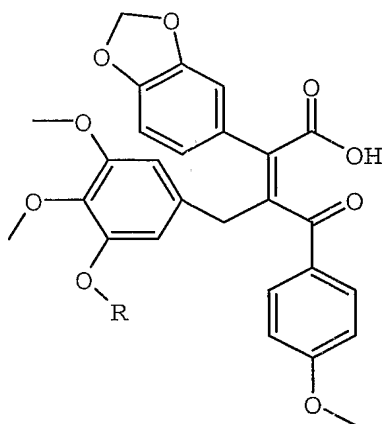
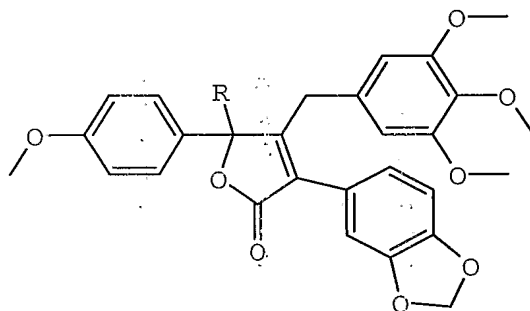
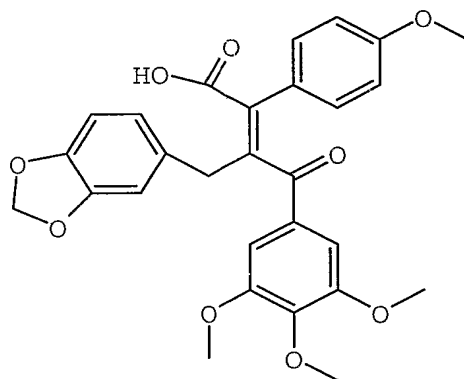
9

BMS 182,874

10 $R_1 = \text{CH}_2\text{OH}$, $R_2 = \text{H}$ 11 $R_1 = \text{H}$, $R_2 = 2\text{-oxazolyl}$ 12 $R_1 = \text{H}$, $R_2 = 2\text{-pyrimidinyl}$ 13 $R_1 = \text{H}$, $R_2 = 4\text{-methoxyethoxymethyl-4-oxo-1,2,4-triazol-2-yl}$ 14 $R_1 = \text{H}$, $R_2 = 1,3\text{-diazol-2-butyl-4-oxospiro(4,4)-1-nonen-3-ylmethyl}$ 

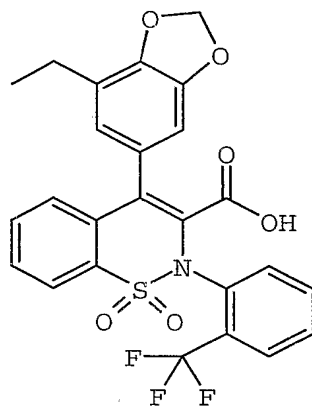
15

- 49 -

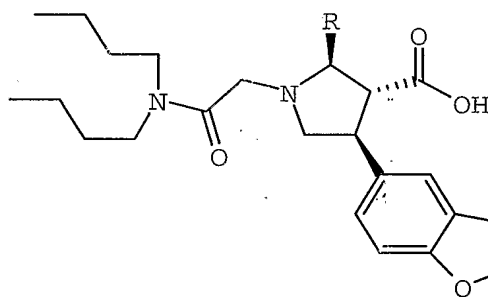
16 $R=CH_3$ (PD156707)17 $R=CH_2CH_2CH_2SO_3H$ 18 $R=OCH_2CH_2CH_2SO_3H$ 19 $R=OCONHCH_2CO_2C_2H_5$ 

20

- 50 -



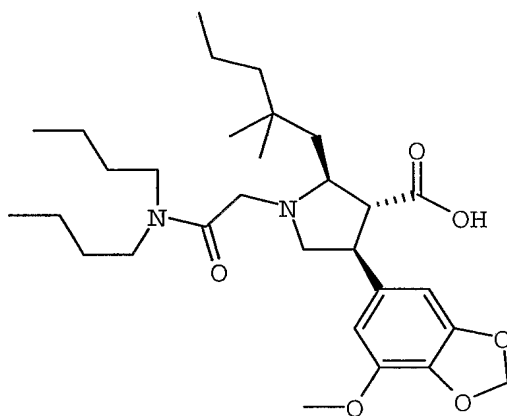
21
PD180988



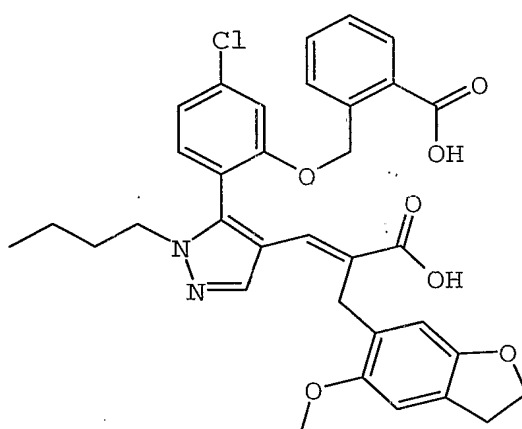
22 R=C₆H₄-4-OCH₃ (ABT-627)

23 R=CH₂CH₂-2-pyridyl

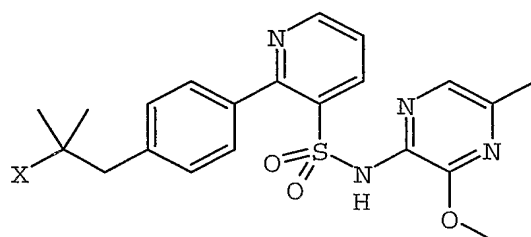
- 51 -



24
ABT-546

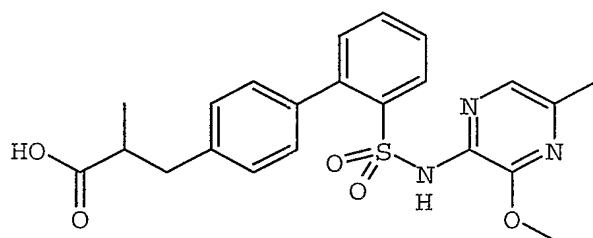


25
SB247083

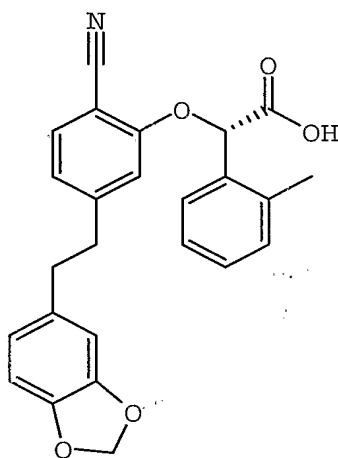


26 X=CO₂H (Z1611)
27 X=H

- 52 -

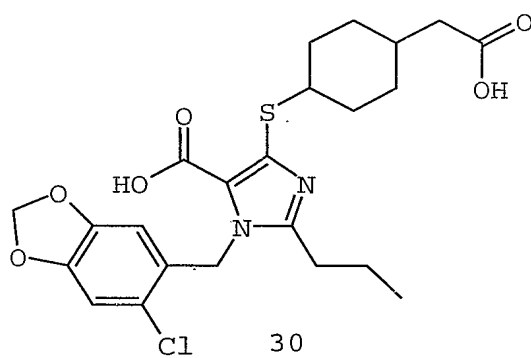


28



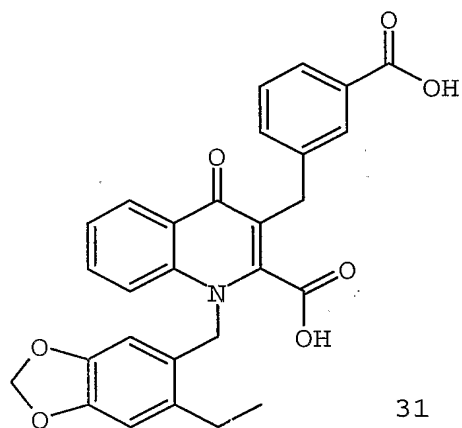
RPR118031A

29

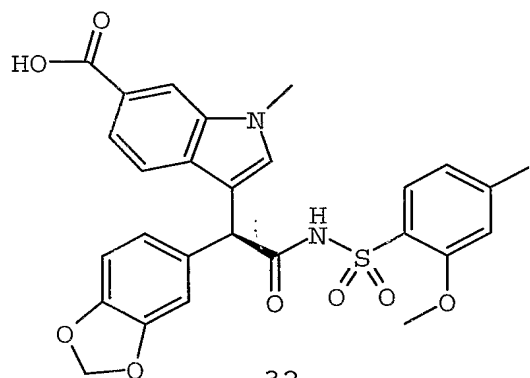


30

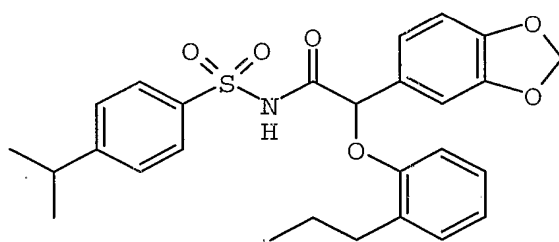
- 53 -



31

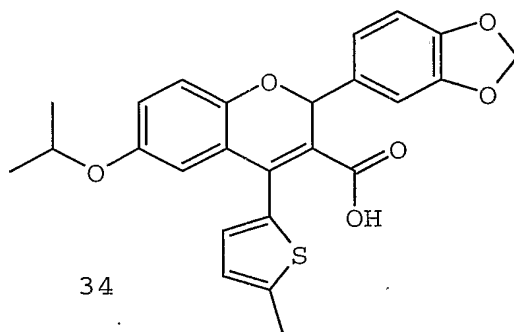


32

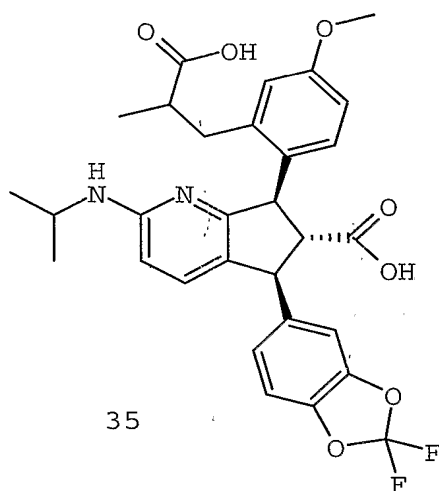


33

- 54 -



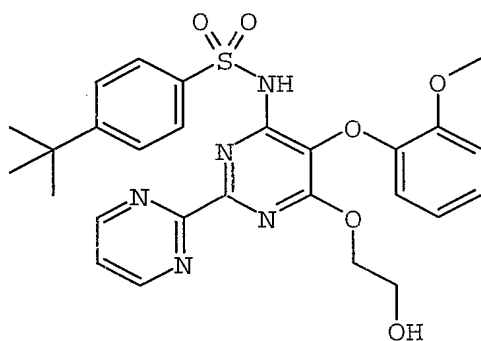
34



35

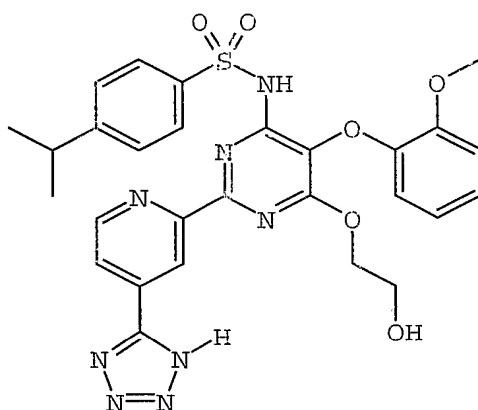
- 55 -

APPENDIX B
BALANCED ET_A/ET_B ANTAGONISTS

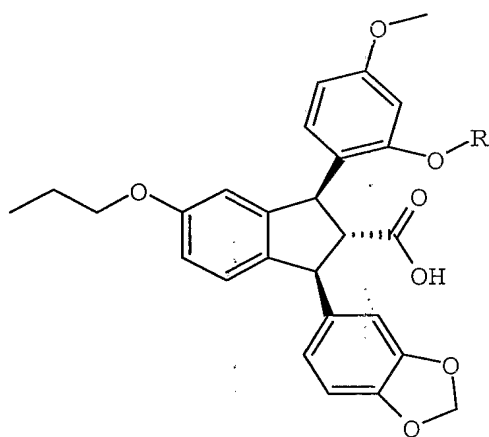


bosentan

46

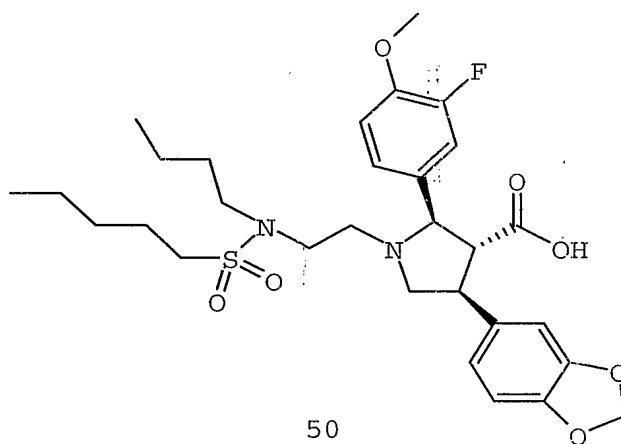


47

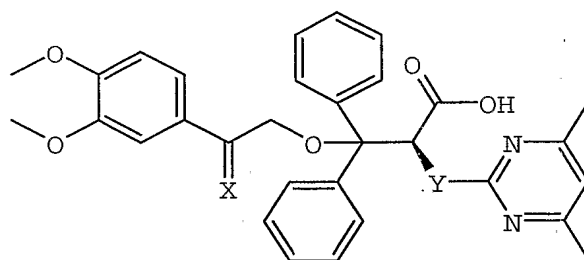


48 R=CH₂CO₂H SB209670

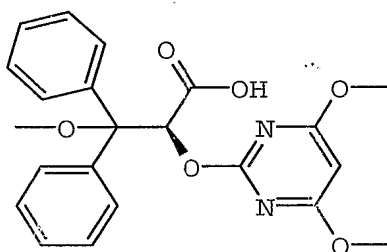
49 R=CH₂CH₂OH SB217242



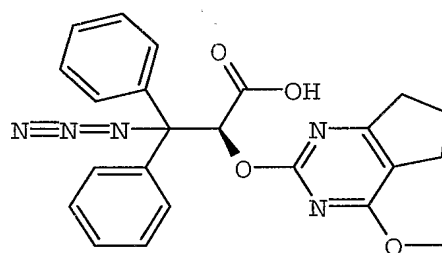
- 57 -

51 X=H₂, Y=CH₂ S-LU 302872

52 X=O, Y=O

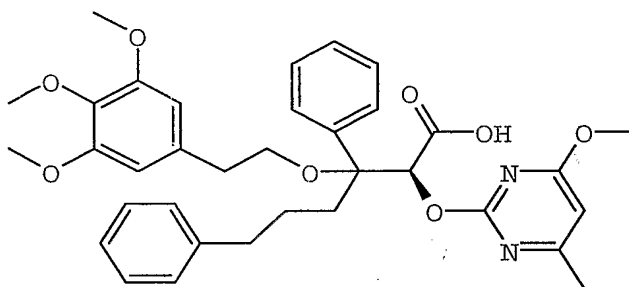


53

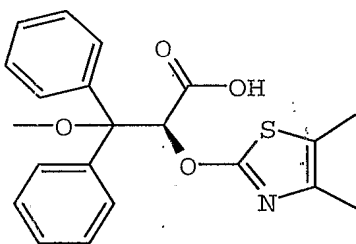


54

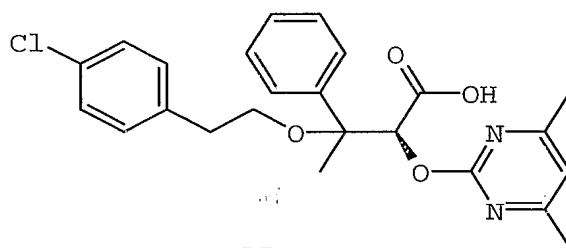
- 58 -



55

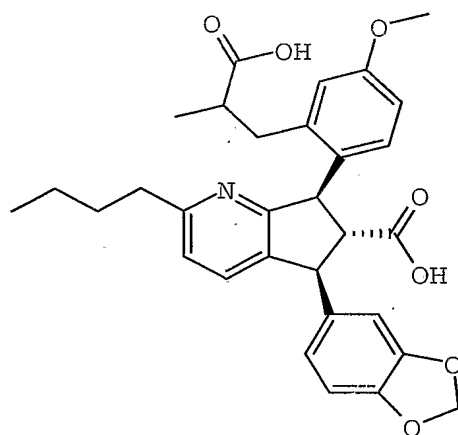


56



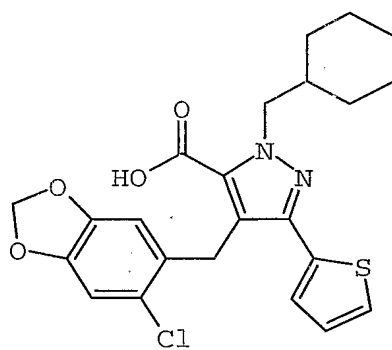
57

- 59 -

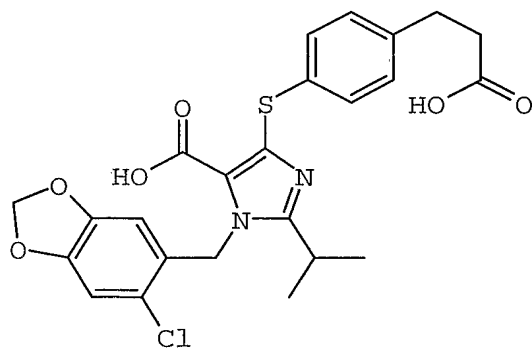


J-104132

58

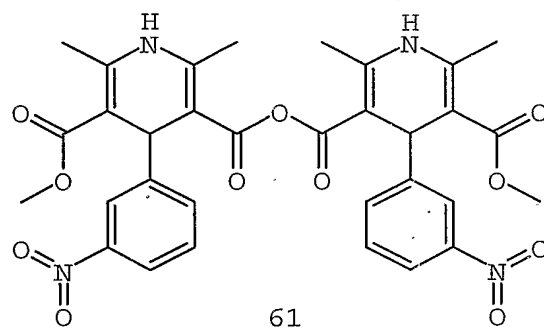


59

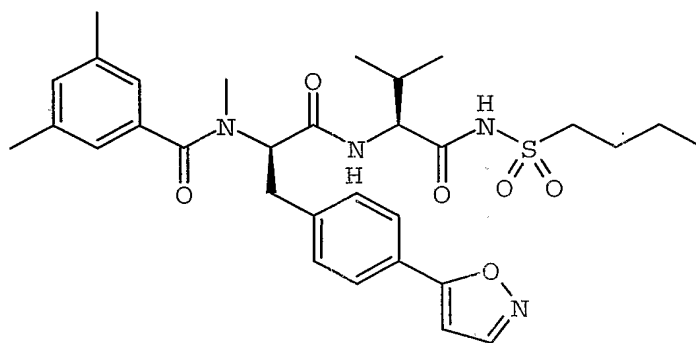


60

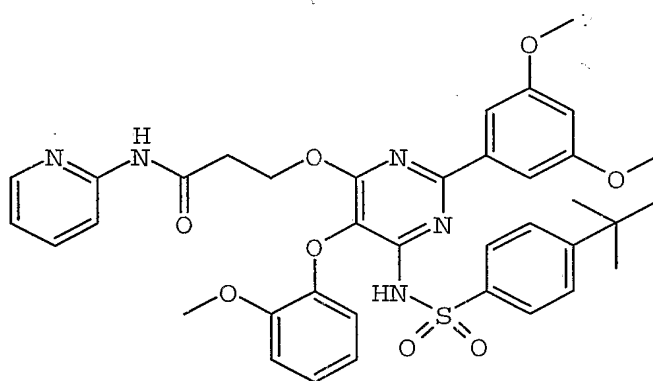
- 60 -



61

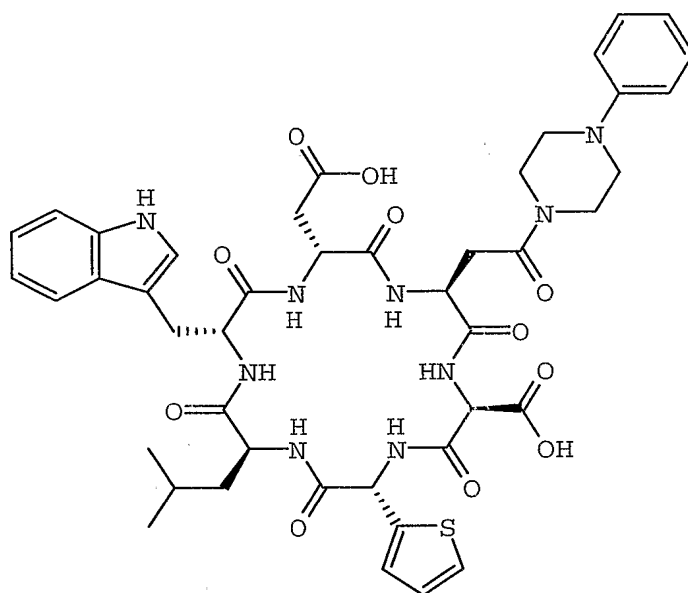


62



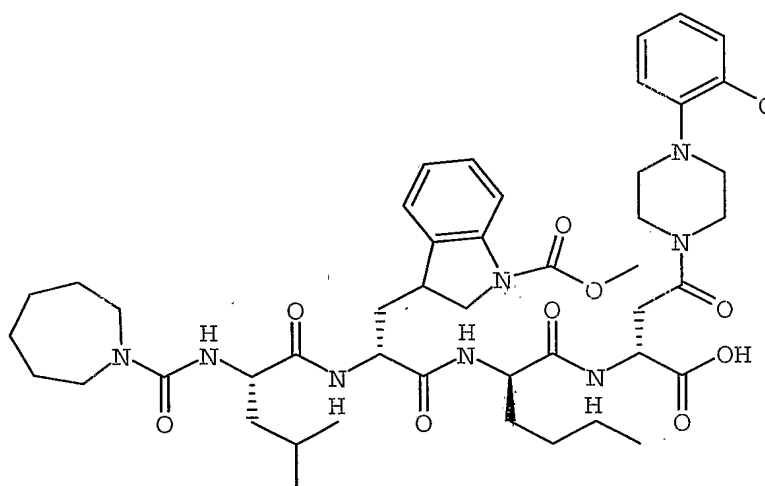
63

- 61 -



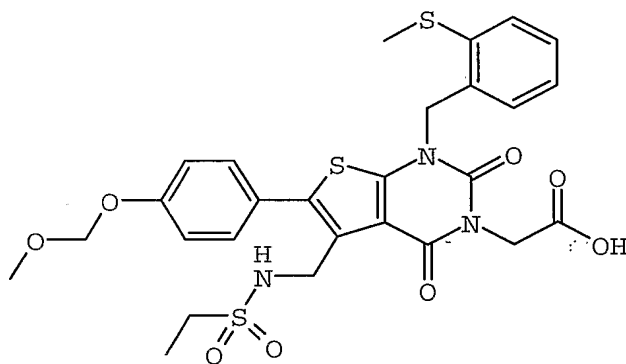
TAK-044

64

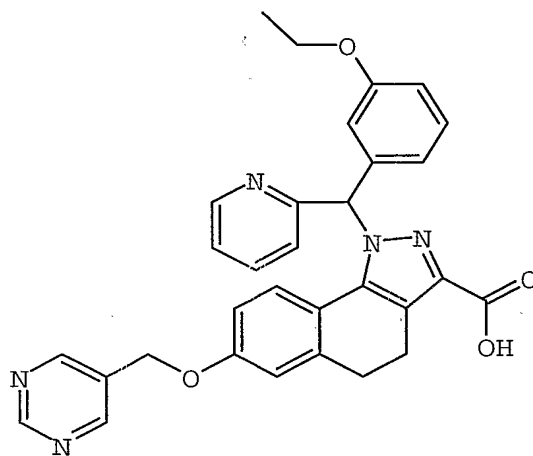


65

- 62 -



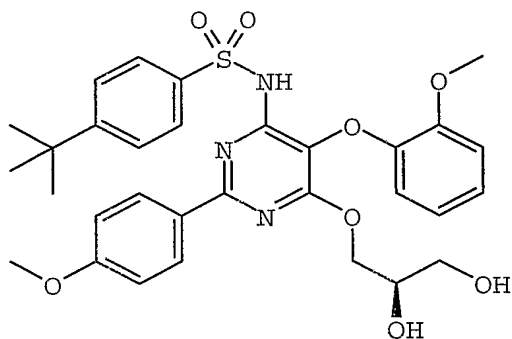
66



67

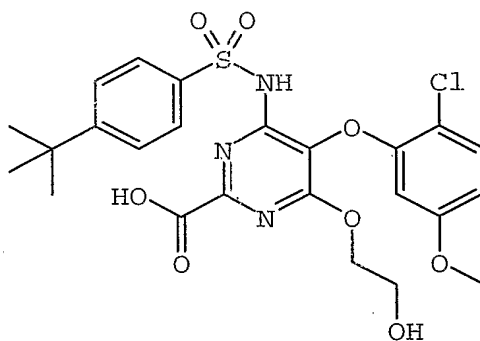
- 63 -

APPENDIX C
SELECTIVE ET_B ANTAGONISTS

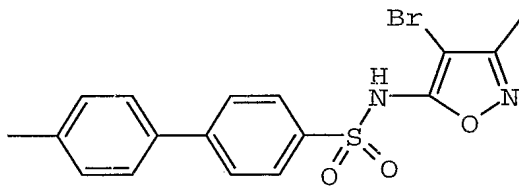


Ro 46-8443

36



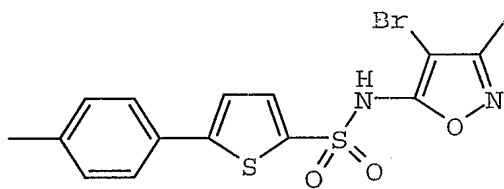
37



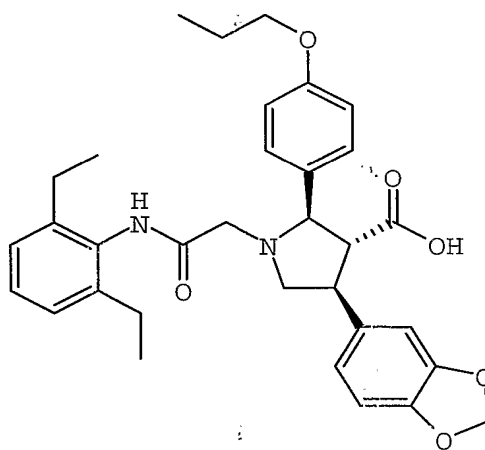
TBC10950

38

- 64 -

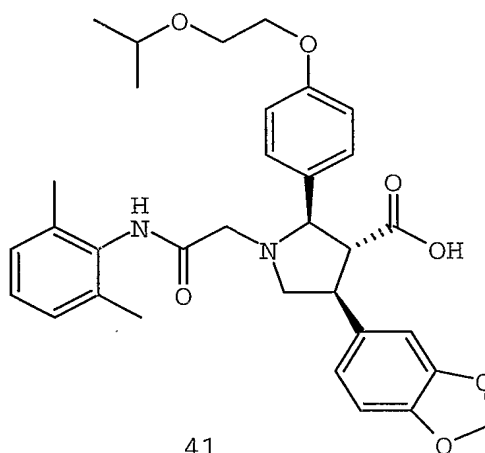


39



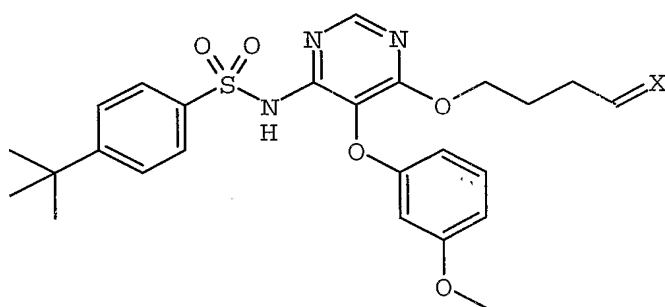
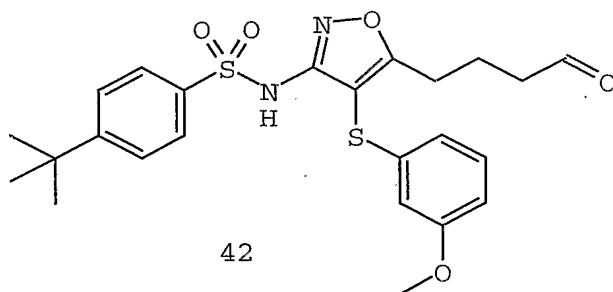
A192621

40

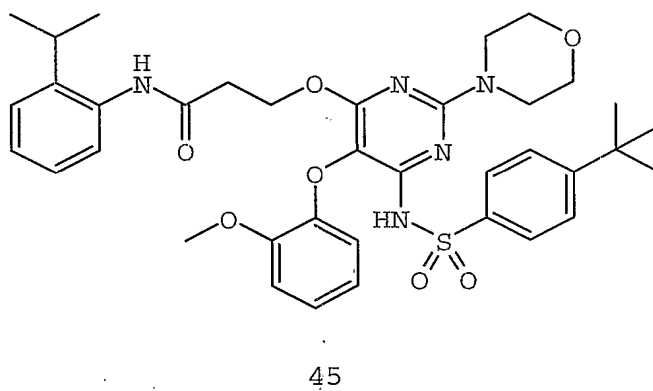


41

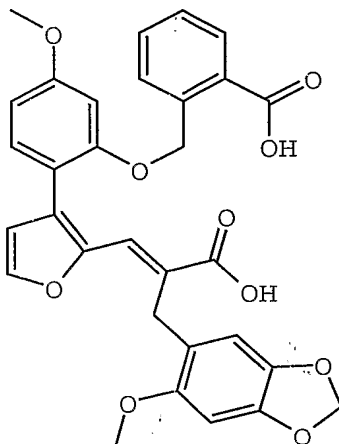
- 65 -



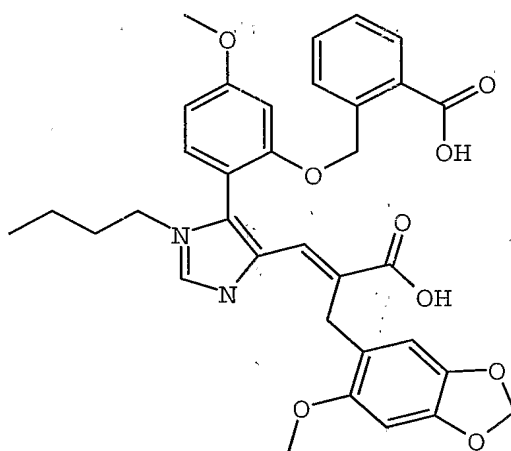
44 X=NNHCO-3-pyridyl



APPENDIX D
MISCELLANEOUS ET ANTAGONISTS

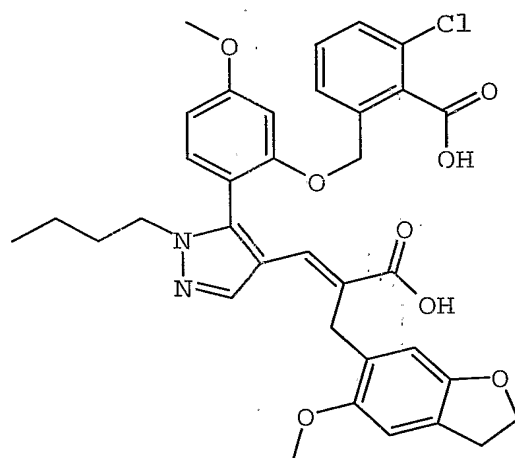
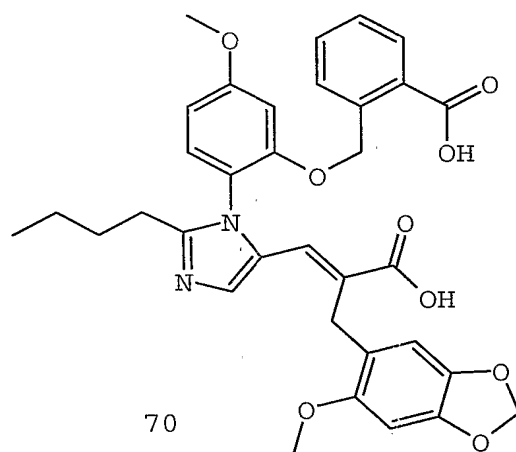


68

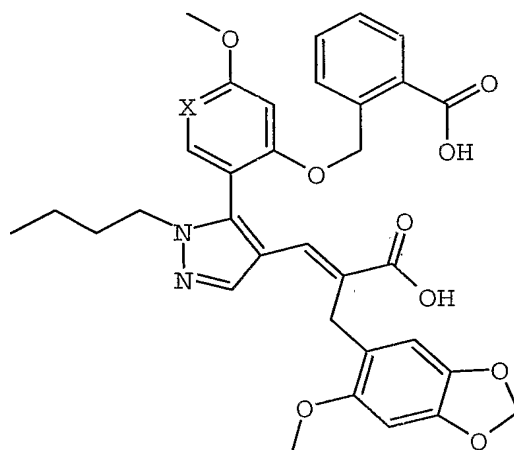


69

- 67 -

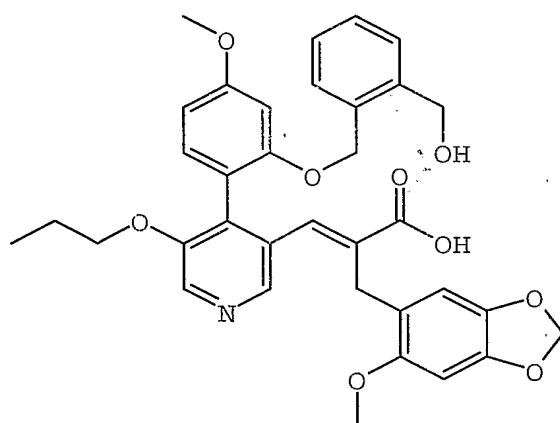


- 68 -



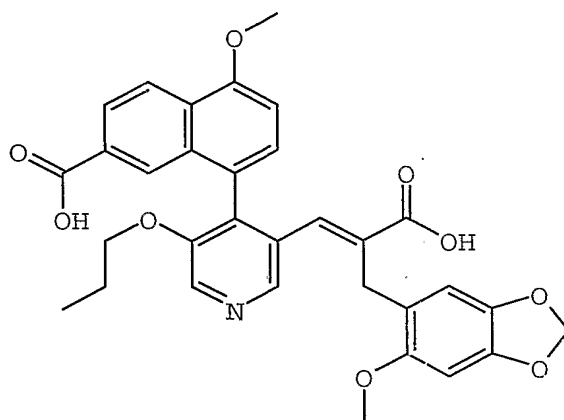
72 X=C

73 X=N

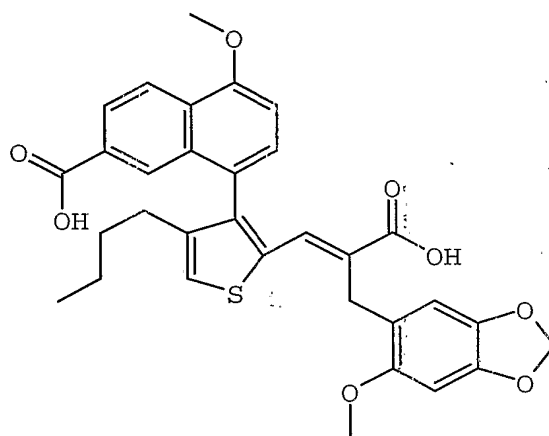


74

- 69 -

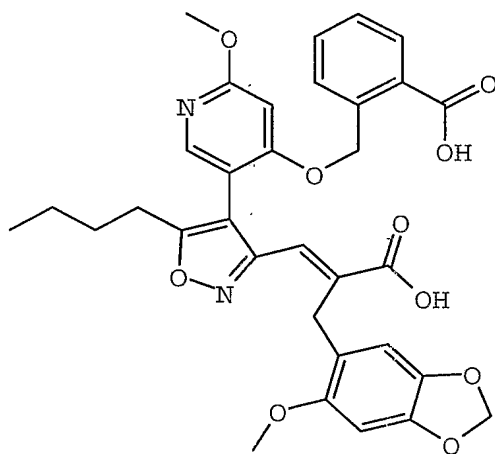


75

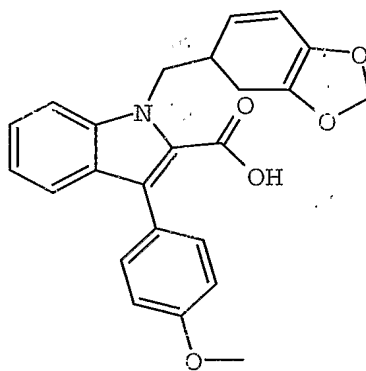


76

- 70 -

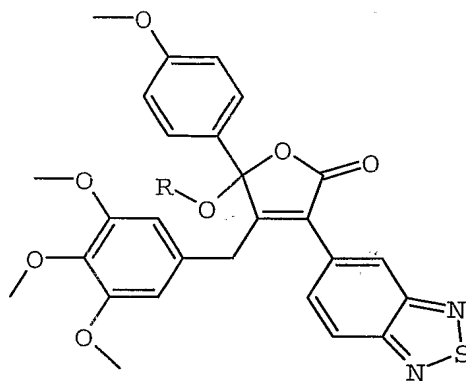


77

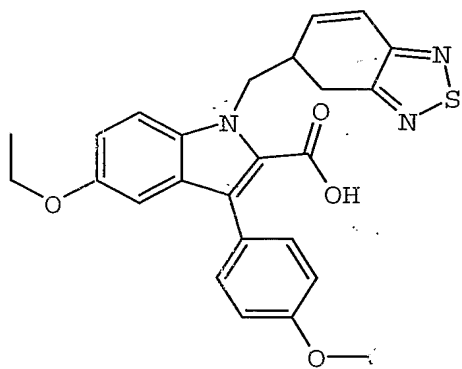


78

- 71 -

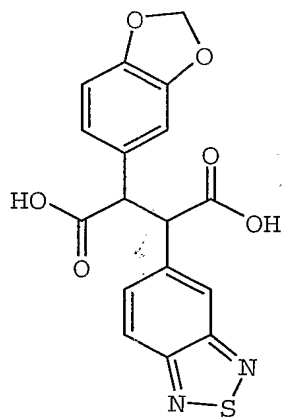


79 R=H

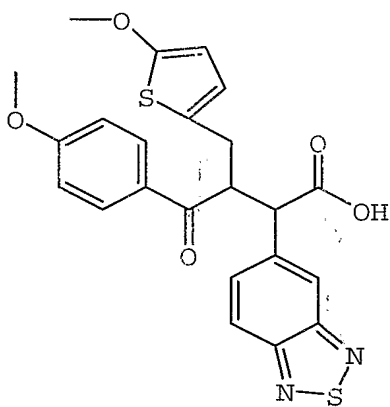
80 R=CONHCH₂CO₂C₂H₅

81

- 72 -

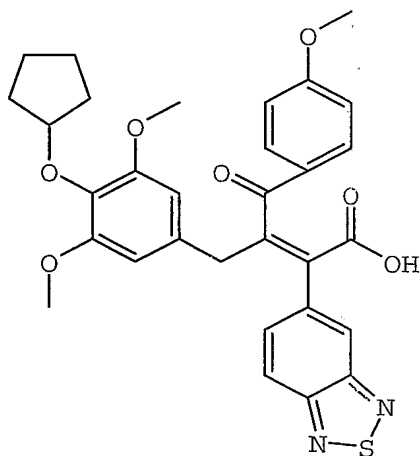


82

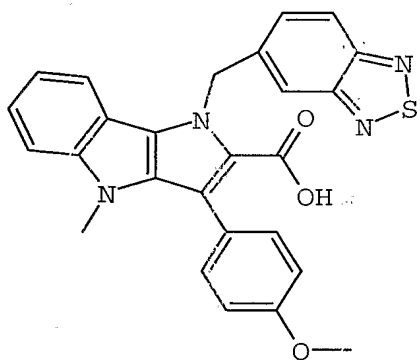


83

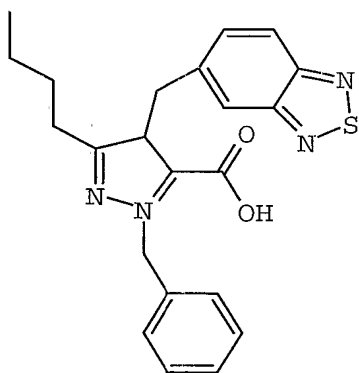
- 73 -



84

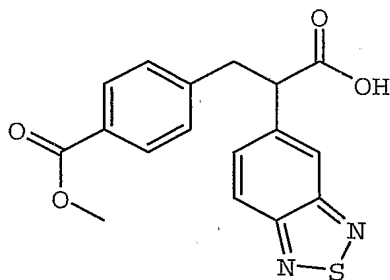


85

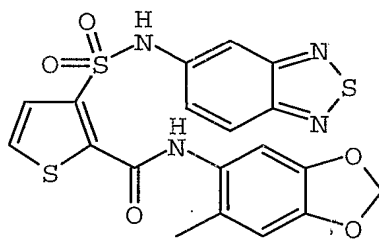


86

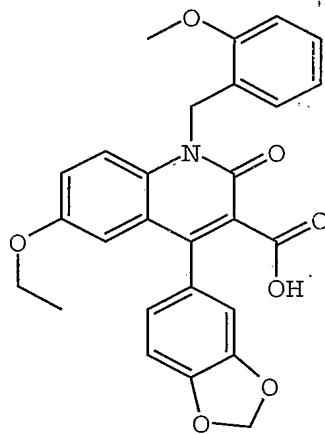
- 74 -



87

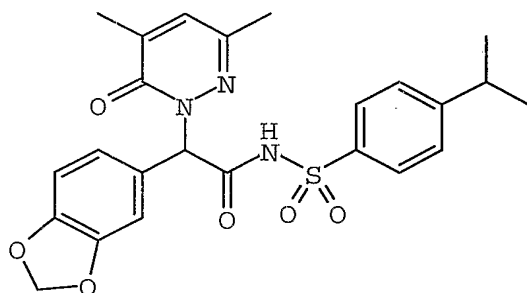


88

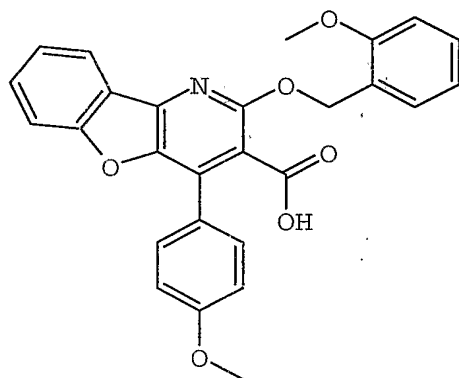


89

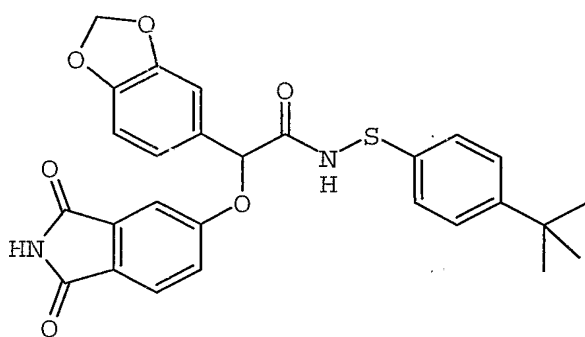
- 75 -



90

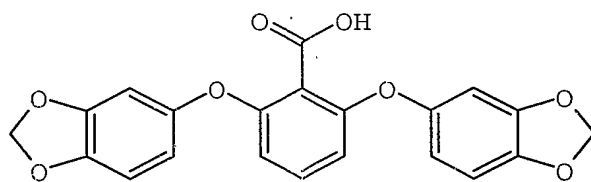


91

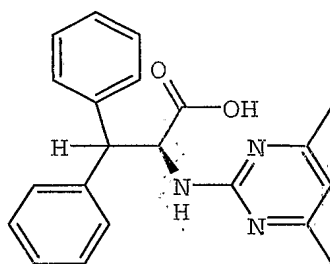


92

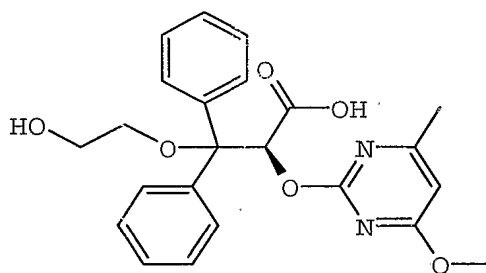
- 76 -



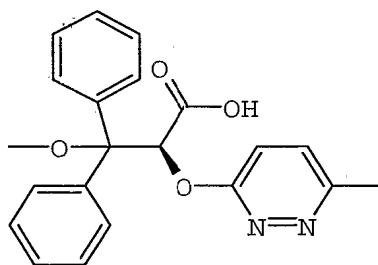
93



94

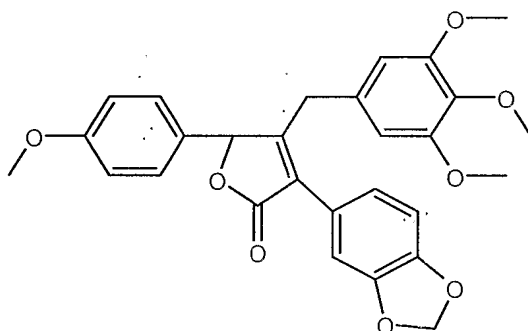


95

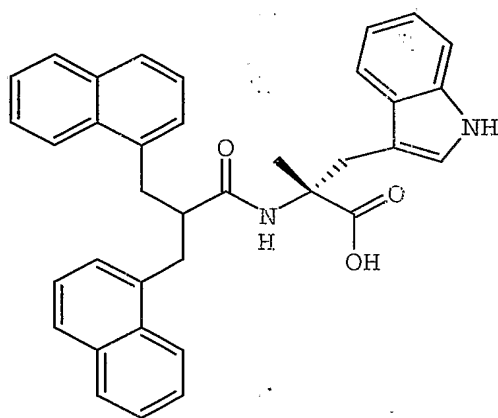


96

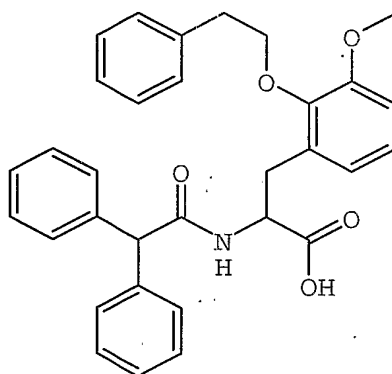
- 77 -



97

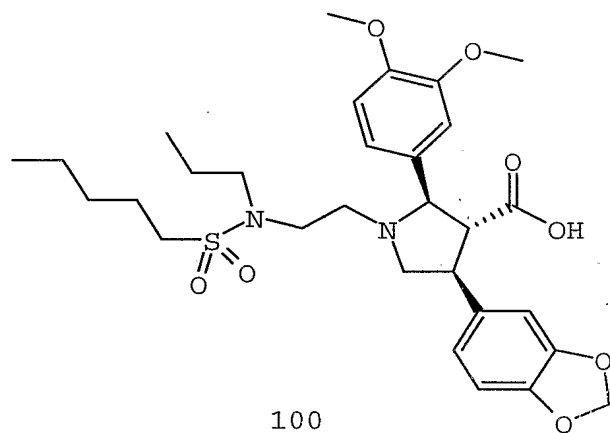


98

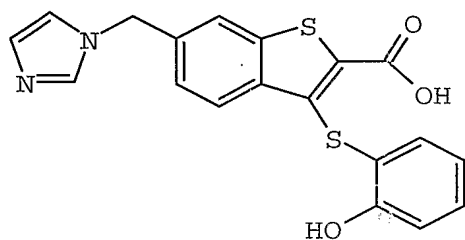


99

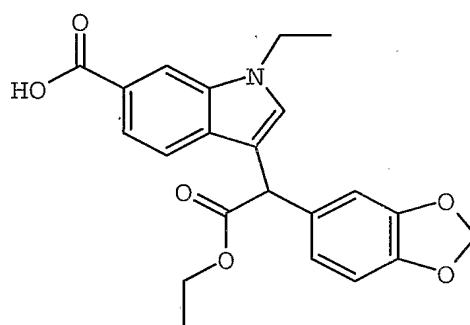
- 78 -



100

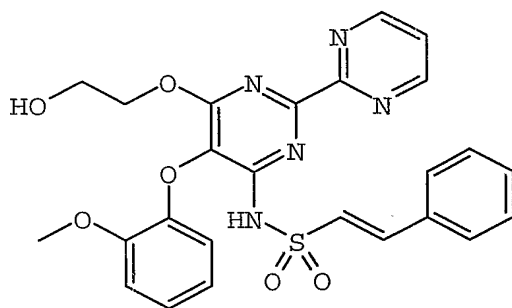


101

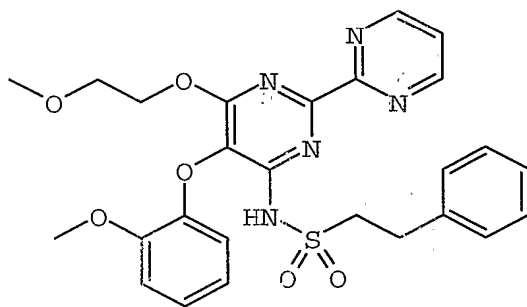


102

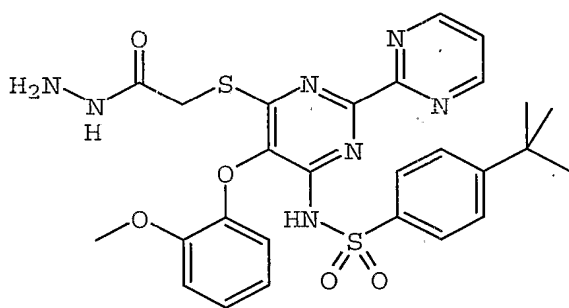
- 79 -



103

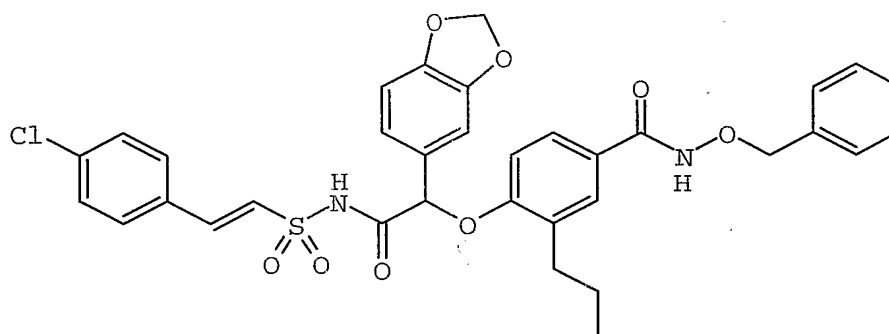


104

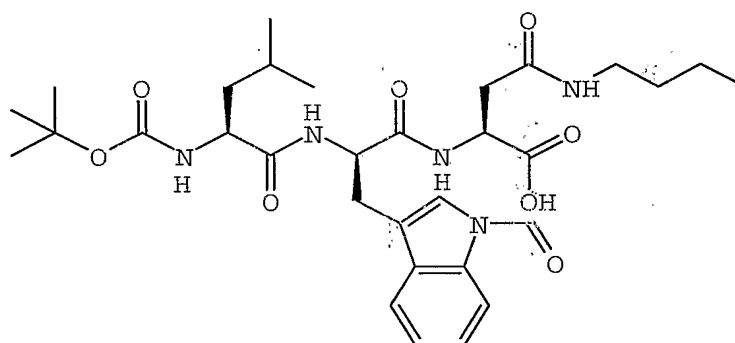


105

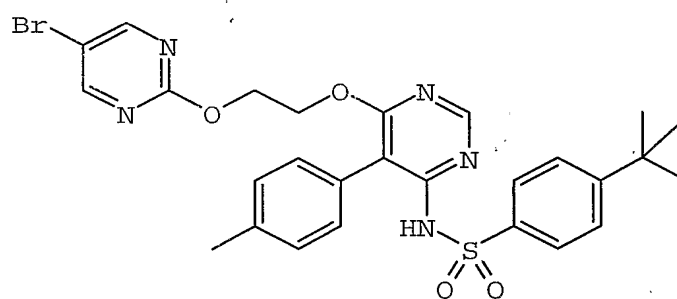
- 80 -



106

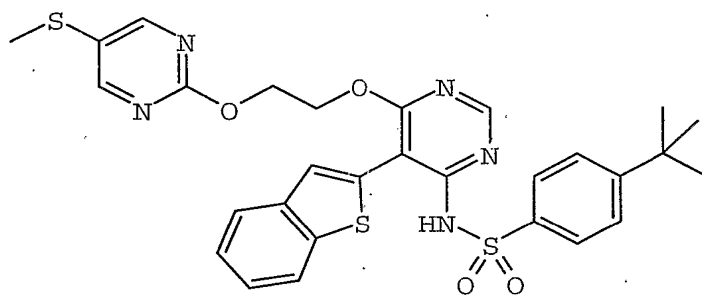


107



108

- 81 -



109

- 82 -

WHAT IS CLAIMED IS:

1. A method of treating or preventing pain comprising administering to a mammal in need thereof a therapeutically effective amount of (a) an opiate analgesic and (b) a therapeutically effective amount of an endothelin antagonist.

2. The method of claim 1 wherein the opiate analgesic and endothelin antagonist are administered simultaneously.

3. The method of claim 2 wherein the opiate analgesic and endothelin antagonist are administered from a single composition.

4. The method of claim 2 wherein the opiate analgesic and endothelin antagonist are administered from separate compositions.

5. The method of claim 1 wherein the opiate analgesic and endothelin antagonist are administered sequentially.

6. The method of claim 5 wherein the opiate analgesic is administered prior to the endothelin antagonist.

- 83 -

7. The method of claim 5 wherein the endothelin antagonist is administered prior to the opiate analgesic.

8. The method of claim 1 wherein the opiate analgesic is selected from the group consisting of an opium alkaloid, a semisynthetic opiate analgesic, and a mixture thereof.

9. The method of claim 1 wherein the opiate analgesic is selected from the group consisting of opium, morphine, morphine sulfate, codeine, codeine phosphate, codeine sulfate, diacetylmorphine, morphine hydrochloride, morphine tartrate, diacetylmorphine hydrochloride, dextromethorphan hydrobromide, hydrocodone bitartrate, hydromorphone, hydromorphone hydrochloride, levorphanol tartrate, oxymorphone hydrochloride, oxycodone hydrochloride, fentanyl, meperidine, methodone, propoxyphene, and mixtures thereof.

10. The method of claim 1 wherein the opiate analgesic comprises morphine or a salt thereof.

- 84 -

11. The method of claim 1 wherein the endothelin antagonist comprises an endothelin-A antagonist.

12. The method of claim 11 wherein the endothelin-A antagonist comprises a specific endothelin-A antagonist.

13. The method of claim 11 wherein the endothelin-A antagonist comprises a nonspecific endothelin-A antagonist.

14. The method of claim 1 wherein the endothelin antagonist is selected from the group consisting of compounds 1 through 35 of Appendix A.

15. The method of claim 1 wherein the endothelin antagonist is selected from the group consisting of compounds 46 through 67 of Appendix B.

16. The method of claim 1 wherein the endothelin antagonist is selected from the group consisting of compounds 36 through 45 of Appendix C.

- 85 -

17. The method of claim 1 wherein the endothelin antagonist is selected from the group consisting of compounds 68 through 109 of Appendix D.

18. The method of claim 1 wherein the endothelin antagonist is selected from the group consisting of atrasentan, tezosentan, bosentan, darnsenten, sitaxsentan, enrasentan, BMS-207940, BMS-193884, BMS-182874, J-104132, VML 588/Ro 61-1790, T-0115, TAK-044, BQ-788, BQ123, YM-598, LU 135252, PD 145065, A-127722, ABT-627, A-192621, A-182086, TBC3711, BSF208075, S-0139, TBC2576, TBC3214, PD156707, PD180988, ABT-546, ABT-627, Z1611, RPR118031A, SB247083, SB217242, S-Lu302872, TPC10950, SB209670, and mixtures thereof.

19. The method of claim 1 wherein the endothelin antagonist comprises BQ123.

20. The method of claim 1 wherein the endothelin antagonist comprises an endothelin converting enzyme inhibitor.

- 86 -

21. The method of claim 20 wherein the endothelin converting enzyme inhibitor is selected from the group consisting of N-((1-((2(S)-(acetylthio)-1-oxopentyl)-amino)-1-cyclopentyl)-carbonyl-S-4-phenylphenyl-alanine methyl ester), phosphoramidon, and mixtures thereof.

22. The method of claim 1 wherein the opiate analgesic is administered at a dose lower than a dose of the same opiate analgesic administered alone to achieve a predetermined reduction of pain.

23. The method of claim 1 wherein cataleptic properties of the opiate analgesic are essentially unaffected.

24. The method of claim 1 wherein the treatment or prevention of pain has a longer duration than a treatment that administers the opiate analgesic alone.

25. The method of claim 1 wherein the mammal is a human.

- 87 -

26. A method of reducing or reversing tolerance to an opiate analgesic in an individual undergoing opiate analgesic therapy comprising administration of a therapeutically effective amount of an endothelin antagonist.

27. A composition comprising (a) an opiate analgesic, (b) an endothelin antagonist, and (c) an optional excipient.

28. The composition of claim 27 wherein the endothelin antagonist comprises an endothelin-A antagonist.

29. An article of manufacture comprising:

- (a) a packaged composition comprising an opiate analgesic;
- (b) a packaged composition comprising an endothelin antagonist;
- (c) an insert providing instructions for a simultaneous or sequential administration of (a) and (b) to treat pain in a mammal; and
- (d) a container for (a), (b), and (c).

- 88 -

30. An article of manufacture comprising:

(a) a packaged composition comprising an opiate analgesic and an endothelin antagonist;

(b) an insert providing instructions for a simultaneous or sequential administration of (a) and (b) to treat pain in a mammal; and

(c) a container for (a) and (b).

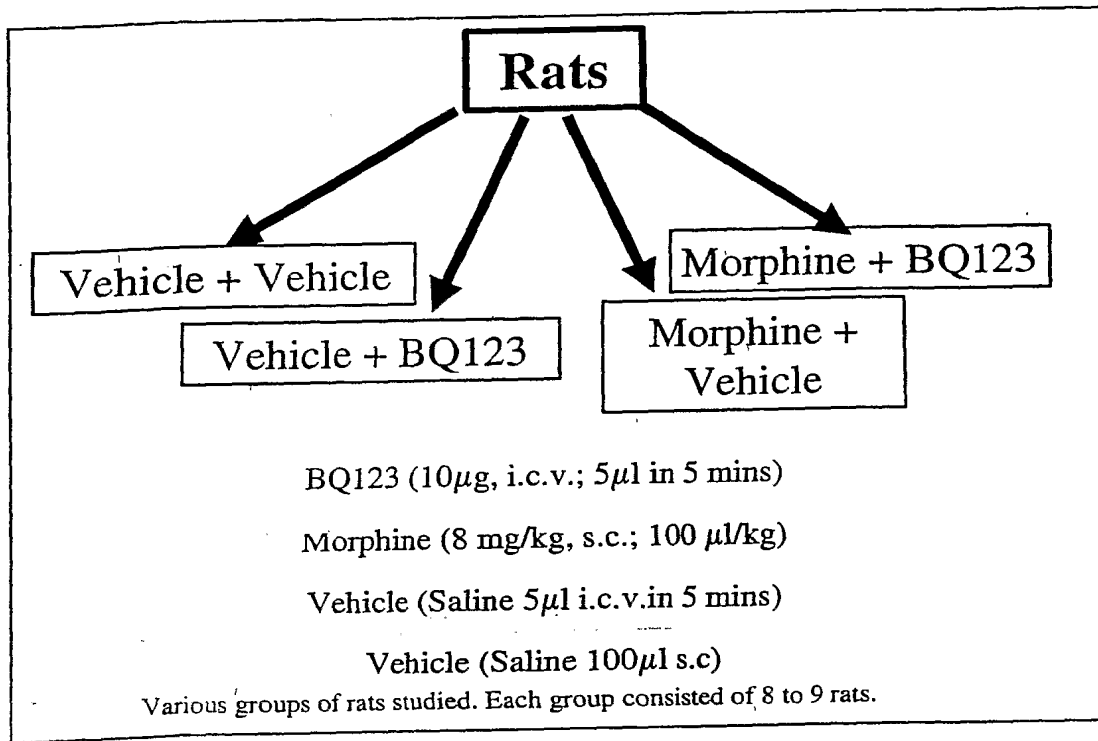
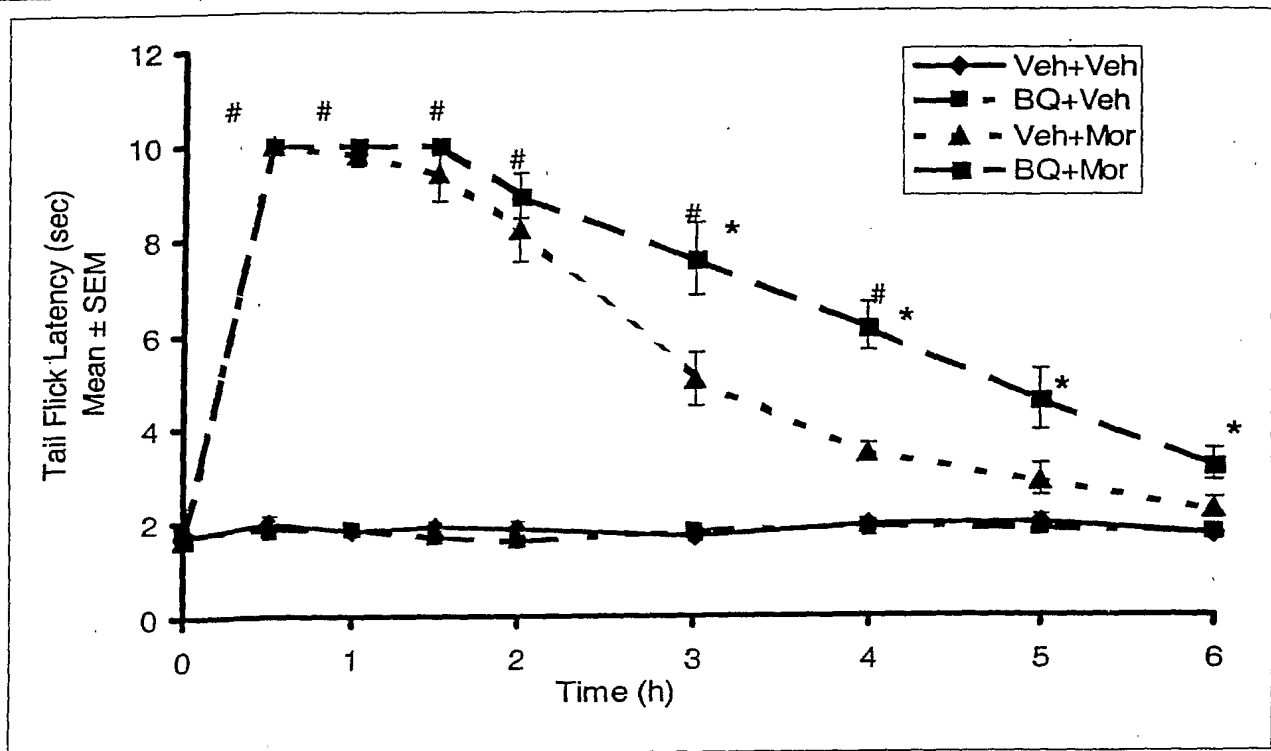
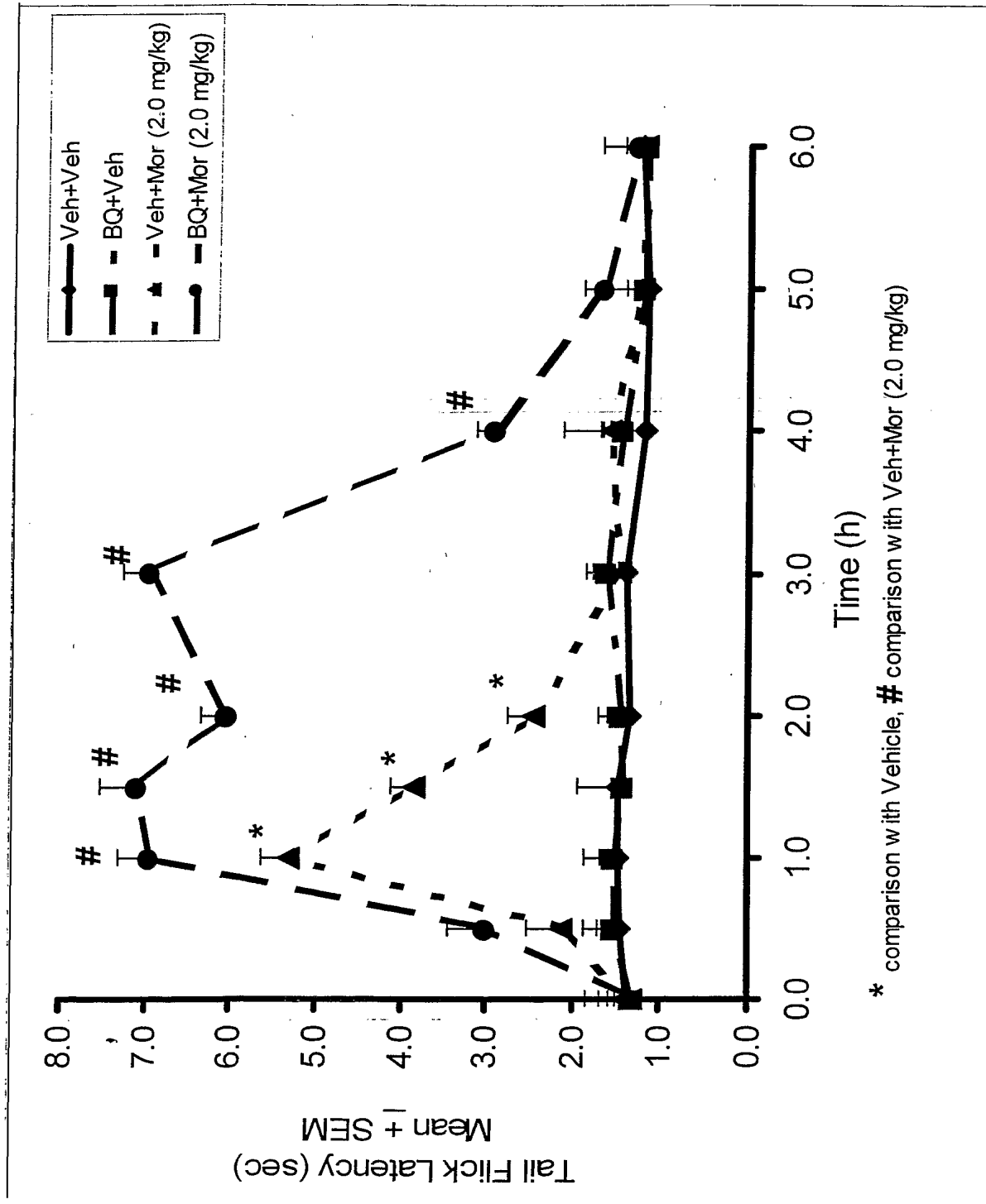


Fig. 1



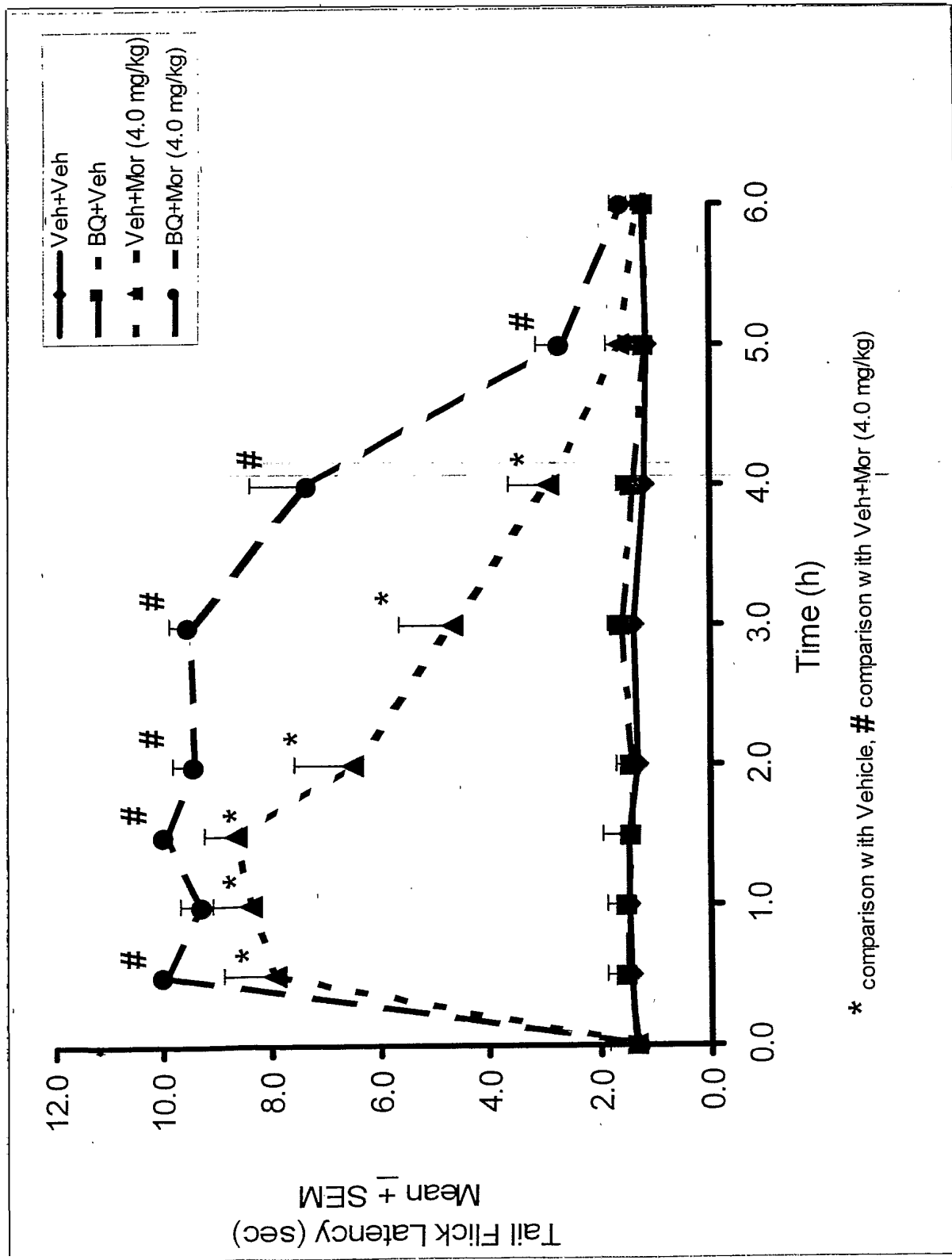
Effect of BQ123 pretreatment on analgesia induced by morphine.

Fig. 2



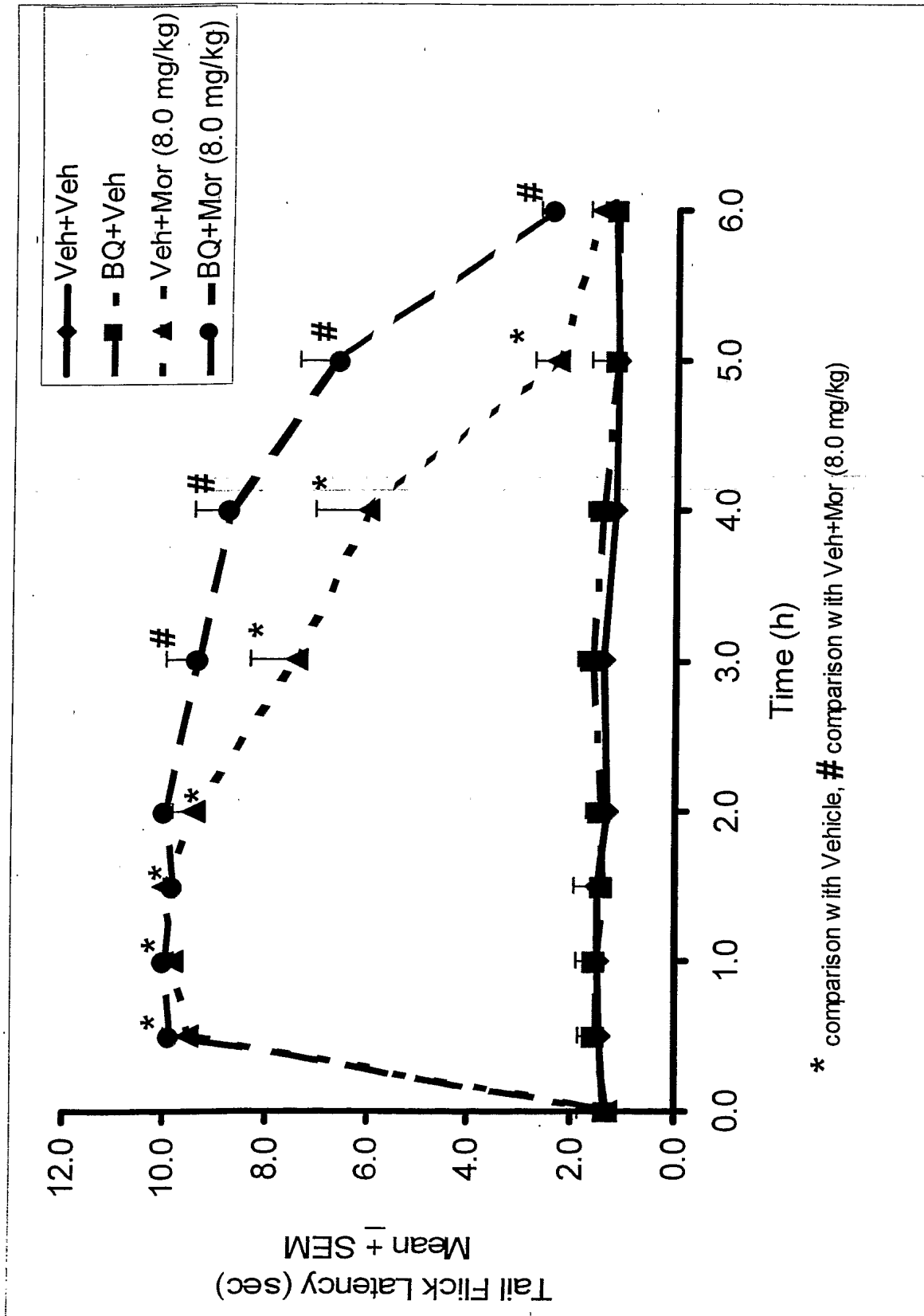
Effect of BQ123 on morphine (2 mg/kg)-induced analgesia

Fig. 3



Effect of BQ123 on morphine (4 mg/kg)-induced analgesia

Fig. 4



Effect of BQ123 on morphine (8 mg/kg)-induced analgesia

Fig. 5

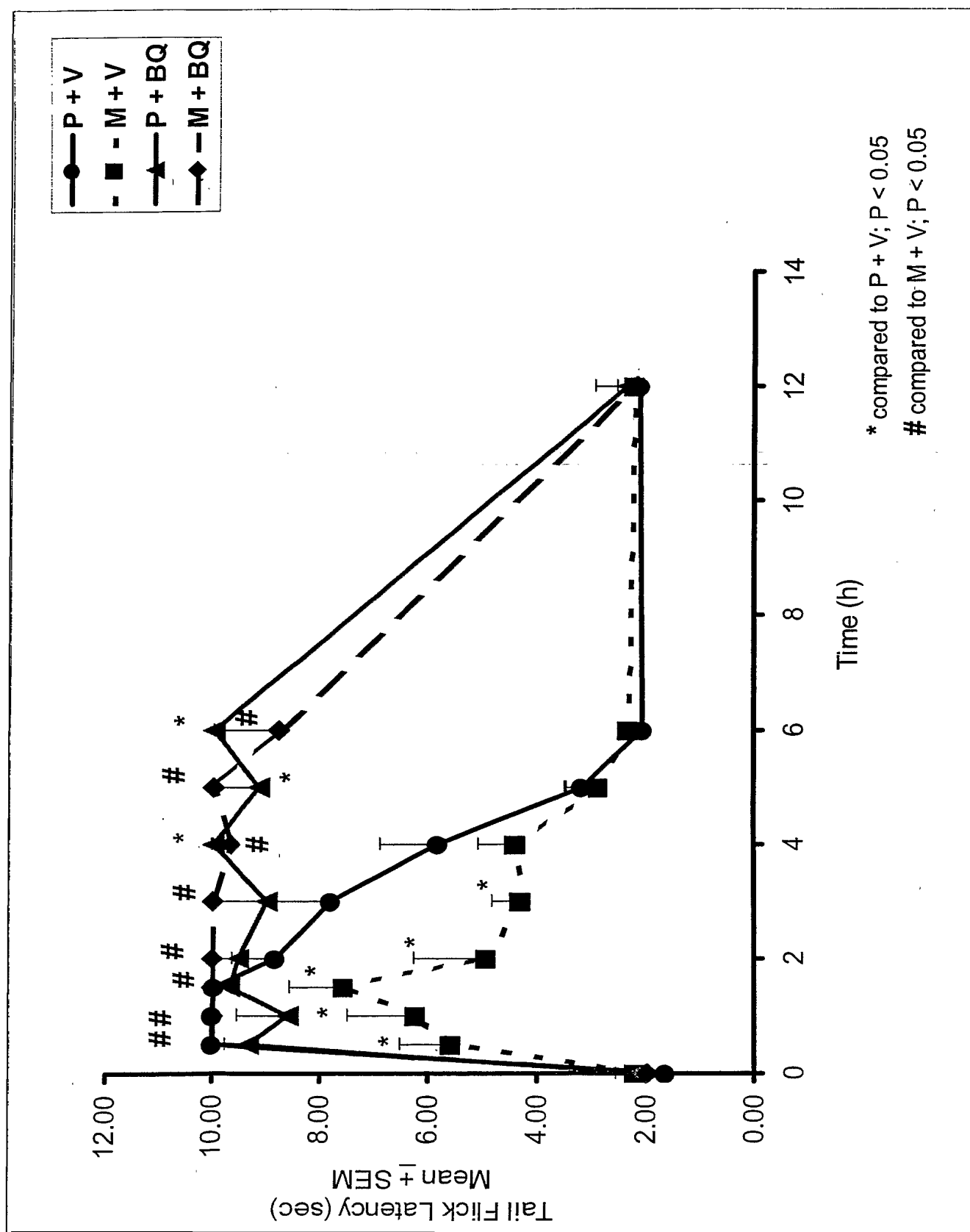
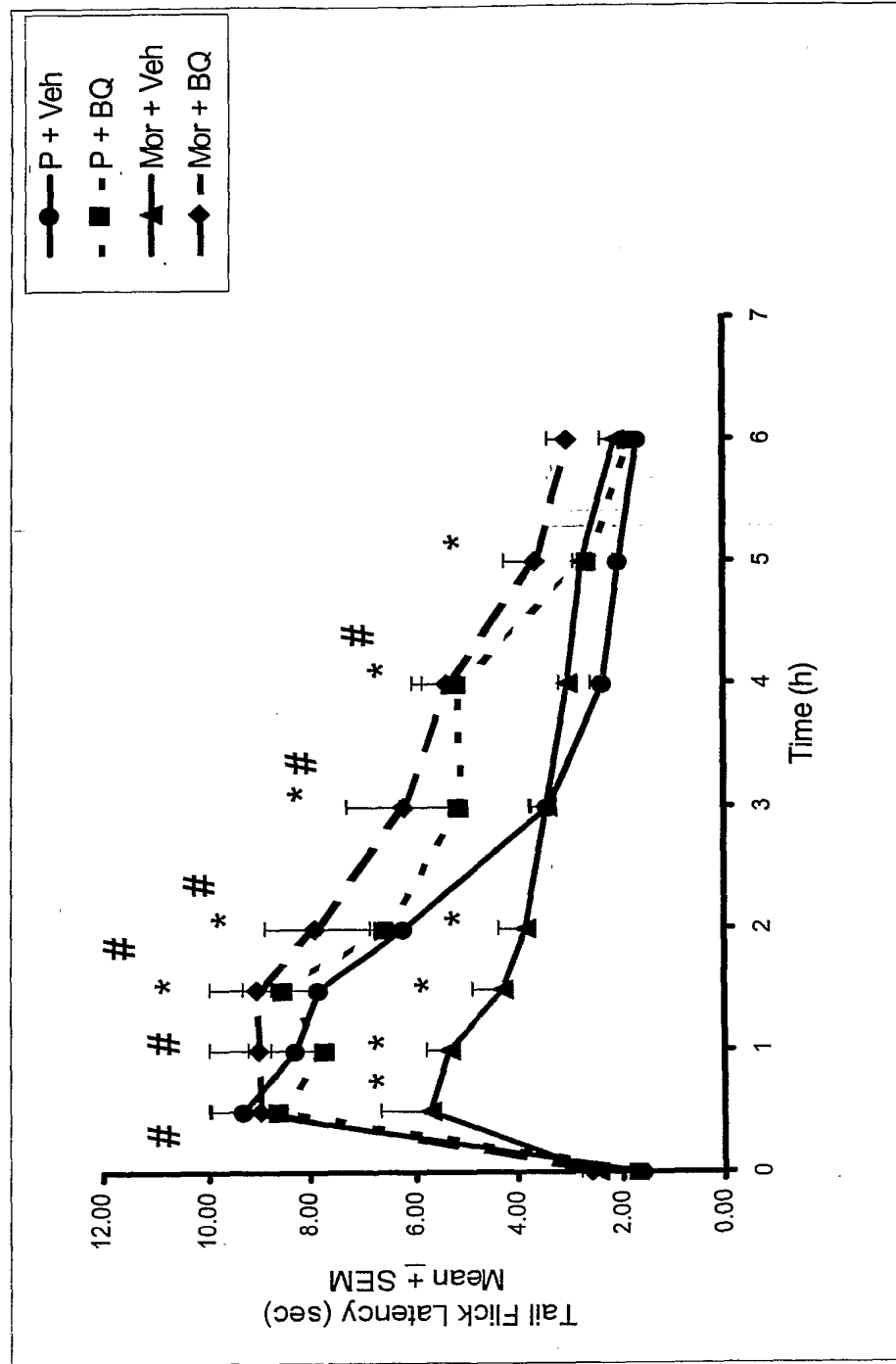


Fig. 6

Effect of BQ123 on morphine induced analgesia in morphine tolerant mice



* p < 0.05 significantly different from control; n=7

p < 0.05 significantly different from morphine; n=7

Fig. 7

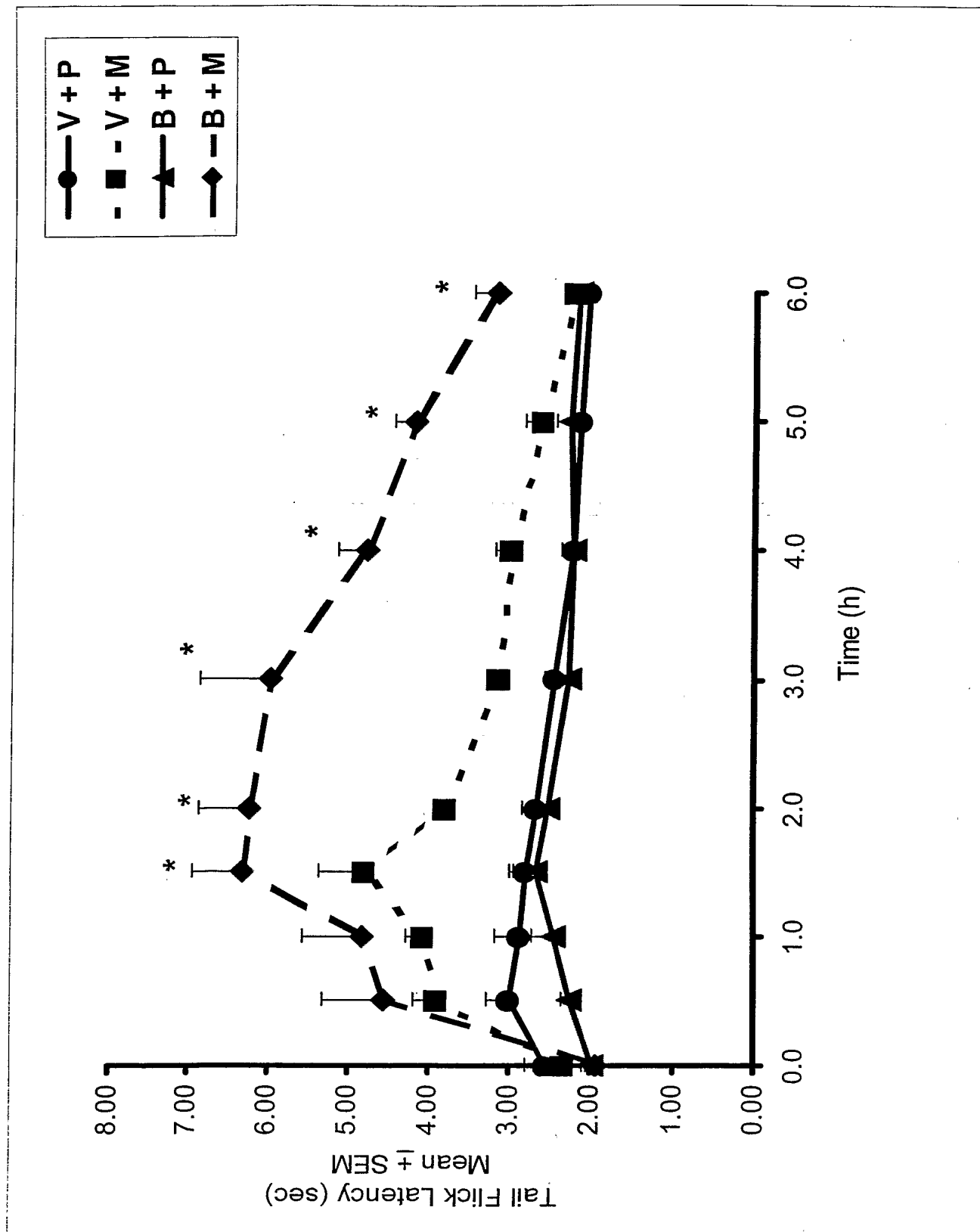


Fig. 8

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
5 June 2003 (05.06.2003)

PCT

(10) International Publication Number
WO 03/045434 A3

(51) International Patent Classification⁷: **A61K 45/06**,
38/12, 31/4025, 31/506 // (A61K 31/422, 31:485), A61P
25/04

(21) International Application Number: PCT/US02/37461

(22) International Filing Date:
22 November 2002 (22.11.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/333,599 27 November 2001 (27.11.2001) US

(71) Applicant: **THE BOARD OF TRUSTEES OF THE
UNIVERSITY OF ILLINOIS** [US/US]; 352 Administra-
tion Building, 506 Wright Street, Urbana, IL 61801 (US).

(72) Inventor: **GULATI, Anil**; 1929 Lisson Road, Naperville,
IL 60565 (US).

(74) Agent: **NAPOLI, James, J.**; Marshall, Gerstein & Borun,
6300 Sears Tower, 233 South Wacker Drive, Chicago, IL
60606 (US).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC,
VN, YU, ZA, ZM, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK,
TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report
- before the expiration of the time limit for amending the
claims and to be republished in the event of receipt of
amendments

(88) Date of publication of the international search report:
25 September 2003

*For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.*

(54) Title: ENDOTHELIN ANTAGONISTS INA METHOD AND COMPOSITION FOR POTENTIATING AN OPIATE ANAL-
GESIC

(57) Abstract: Composition and methods of treating pain and reducing or reversing tolerance to opiate analgesics are disclosed. The
composition and method utilize an opiate analgesic and an endothelin antagonist as active agents to treat pain in mammals, including
humans.



WO 03/045434 A3

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 02/37461

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K45/06 A61K38/12 A61K31/4025 A61K31/506 A61K31/422
A61K31/485 A61P25/04

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, BIOSIS, MEDLINE, EMBASE, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>WO 96 19233 A (DEMOPULOS GREGORY A ;HERZ JEFFREY M (US); OMEROS MED SYS INC (US);) 27 June 1996 (1996-06-27) page 9, line 19 page 24, line 25 page 33, line 1-22; table 20</p> <p style="text-align: center;">----- -/--</p>	<p>1-25, 29, 30</p>

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

31 July 2003

Date of mailing of the international search report

06/08/2003

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Baumgärtner, H

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 02/37461

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	JARVIS M F ET AL: "ABT-627, AN ENDOTHELIN ETA RECEPTOR-SELECTIVE ANTAGONIST, ATTENUATES TACTILE ALLODYNIA IN A DIABETIC RAT MODEL OF NEUROPATHIC PAIN" EUROPEAN JOURNAL OF PHARMACOLOGY, AMSTERDAM, NL, vol. 388, no. 1, 24 January 2000 (2000-01-24), pages 29-35, XP001152675 ISSN: 0014-2999 page 30, column 1, paragraph 3 page 31, column 2, last paragraph -page 32, column 1, paragraph 1 page 31; figure 2 ----	1-30
Y	DAVAR G ET AL: "BEHAVIORAL SIGNS OF ACUTE PAIN PRODUCED BY APPLICATION OF ENDOTHELIN-1 TO RAT SCIATIC NERVE" NEUROREPORT, RAPID COMMUNICATIONS OF OXFORD, OXFORD, GB, vol. 9, no. 10, 13 July 1998 (1998-07-13), pages 2279-2283, XP008007945 ISSN: 0959-4965 page 2282, column 1, paragraph 1 page 2282, column 2, paragraph 2 page 2282, column 2, last paragraph ----	1-30
A	US 5 612 359 A (MURUGESAN NATESAN) 18 March 1997 (1997-03-18) column 1, line 13 and formula I column 6, line 3-5 column 5, line 49-52 ----	1-30
P,X	BHALLA SHAI FALI ET AL: "Potentiation of morphine analgesia by BQ123, an endothelin antagonist." PEPTIDES (NEW YORK), vol. 23, no. 10, 20 October 2002 (2002-10-20), pages 1837-1845, XP001152674 ISSN: 0196-9781 page 1838, column 2, paragraph 2 page 1844, column 1, paragraph 2 page 1844 sentence bridging column 1-2 -----	1-30

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 02/37461

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

Although claims 1-28 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☒ Claims Nos.: 1, 8, 11-17, 20, 22-24, 27, 29
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

see FURTHER INFORMATION sheet PCT/ISA/210
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1, 8, 11-17, 20, 22-24, 27, 29

Present claims 1, 8, 11-17, 27, 29, 30 relate to an extremely large number of possible compounds. Support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT is to be found, however, for only a very small proportion of the compounds claimed. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Consequently, the search has been carried out for those parts of the claims which appear to be supported and disclosed, namely those parts relating to the compounds as defined in

claims 9/10 (opiate analgesics)
claims 18/19 (endothelin antagonists)
claim 21 (ECE)

Present claims 22-24 relate to features defined by reference to a desirable characteristic or property, namely "lower dose than a dose of the same opiate analgesic administered alone ..." , "wherein the cataleptic properties of the opiate analgesic are essentially unaffected", "has longer duration than a treatment that administers the opiate analgesic alone".

The claims cover all features having this characteristic or property, whereas the application provides support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT for only a very limited number of such features. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Independent of the above reasoning, the claims also lack clarity (Article 6 PCT). An attempt is made to define the features by reference to a result to be achieved. Again, this lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible. Consequently, the search has been carried out for those parts of the claims which appear to be clear, supported and disclosed.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/US 02/37461

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
WO 9619233	A	27-06-1996	CA	2206119 A1	27-06-1996
			CN	1175213 A	04-03-1998
			RU	2180852 C2	27-03-2002
			WO	9619233 A2	27-06-1996
			US	5800385 A	01-09-1998
			US	5860950 A	19-01-1999
			US	5858017 A	12-01-1999
<hr/>					
US 5612359	A	18-03-1997	US	5827869 A	27-10-1998
			AU	699138 B2	26-11-1998
			AU	3026195 A	07-03-1996
			CA	2155447 A1	27-02-1996
			CN	1128262 A ,B	07-08-1996
			CZ	9502163 A3	13-03-1996
			EP	0702012 A1	20-03-1996
			FI	954008 A	27-02-1996
			HU	74097 A2	28-11-1996
			IL	114829 A	30-11-1999
			JP	8183786 A	16-07-1996
			NO	953361 A	27-02-1996
			NZ	272812 A	24-11-1997
			PL	310191 A1	04-03-1996
			RU	2174979 C2	20-10-2001
			SG	46142 A1	20-02-1998
			TW	461890 B	01-11-2001
			BR	1100659 A3	13-06-2000
<hr/>					